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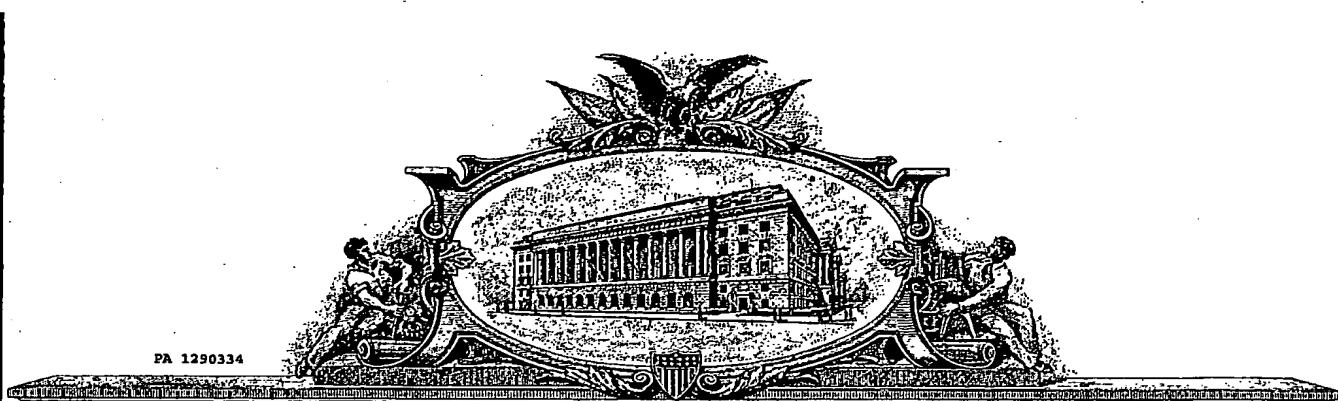
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This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c).33431 U 3296  
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TITLE OF THE INVENTION (280 characters max)	
PHARMACEUTICAL COMPOUNDS	

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Respectfully submitted,

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Date 12/23/03

REGISTRATION NO. 32,700  
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## PHARMACEUTICAL COMPOUNDS

This invention relates to pyrazole-containing aryl- and heteroaryl-alkylamine compounds that inhibit or modulate the activity of protein kinase A (PKA) and 5 protein kinase B (PKB), to the use of the compounds in the treatment or prophylaxis of disease states or conditions mediated by PKA and PKB, and to novel compounds having PKA and PKB inhibitory or modulating activity. Also provided are pharmaceutical compositions containing the compounds and novel chemical intermediates.

10 **Background of the Invention**

Protein kinases constitute a large family of structurally related enzymes that are responsible for the control of a wide variety of signal transduction processes within the cell (Hardie, G. and Hanks, S. (1995) *The Protein Kinase Facts Book. I and II*, Academic Press, San Diego, CA). The kinases may be categorized into families by 15 the substrates they phosphorylate (e.g., protein-tyrosine, protein-serine/threonine, lipids, etc.). Sequence motifs have been identified that generally correspond to each of these kinase families (e.g., Hanks, S.K., Hunter, T., *FASEB J.*, 9:576-596 (1995); Knighton, *et al.*, *Science*, 253:407-414 (1991); Hiles, *et al.*, *Cell*, 70:419-429 (1992); Kunz, *et al.*, *Cell*, 73:585-596 (1993); Garcia-Bustos, *et al.*, *EMBO J.*, 20: 13:2352-2361 (1994)).

Protein kinases may be characterized by their regulation mechanisms. These mechanisms include, for example, autophosphorylation, transphosphorylation by other kinases, protein-protein interactions, protein-lipid interactions, and protein-polynucleotide interactions. An individual protein kinase may be regulated by more 25 than one mechanism.

Kinases regulate many different cell processes including, but not limited to, proliferation, differentiation, apoptosis, motility, transcription, translation and other signalling processes, by adding phosphate groups to target proteins. These phosphorylation events act as molecular on/off switches that can modulate or

regulate the target protein biological function. Phosphorylation of target proteins occurs in response to a variety of extracellular signals (hormones, neurotransmitters, growth and differentiation factors, etc.), cell cycle events, environmental or nutritional stresses, etc. The appropriate protein kinase functions

5 in signalling pathways to activate or inactivate (either directly or indirectly), for example, a metabolic enzyme, regulatory protein, receptor, cytoskeletal protein, ion channel or pump, or transcription factor. Uncontrolled signalling due to defective control of protein phosphorylation has been implicated in a number of diseases, including, for example, inflammation, cancer, allergy/asthma, diseases and

10 conditions of the immune system, diseases and conditions of the central nervous system, and angiogenesis.

Apoptosis or programmed cell death is an important physiological process which removes cells no longer required by an organism. The process is important in early embryonic growth and development allowing the non-necrotic controlled

15 breakdown, removal and recovery of cellular components. The removal of cells by apoptosis is also important in the maintenance of chromosomal and genomic integrity of growing cell populations. There are several known checkpoints in the cell growth cycle at which DNA damage and genomic integrity are carefully monitored. The response to the detection of anomalies at such checkpoints is to

20 arrest the growth of such cells and initiate repair processes. If the damage or anomalies cannot be repaired then apoptosis is initiated by the damaged cell in order to prevent the propagation of faults and errors. Cancerous cells consistently contain numerous mutations, errors or rearrangements in their chromosomal DNA. It is widely believed that this occurs in part because the majority of tumours have a

25 defect in one or more of the processes responsible for initiation of the apoptotic process. Normal control mechanisms cannot kill the cancerous cells and the chromosomal or DNA coding errors continue to be propagated. As a consequence restoring these pro-apoptotic signals or suppressing unregulated survival signals is an attractive means of treating cancer.

The signal transduction pathway containing the enzymes phosphatidylinositol 3-kinase (PI3K), PDK1 and PKB amongst others, has long been known to mediate increased resistance to apoptosis or survival responses in many cells. There is a substantial amount of data to indicate that this pathway is an important survival

5 pathway used by many growth factors to suppress apoptosis. The enzyme PI3K is activated by a range of growth and survival factors e.g. EGF, PDGF and through the generation of polyphosphatidylinositols, initiates the activation of the downstream signalling events including the activity of the kinases PDK1 and protein kinase B (PKB) also known as akt. PKB is a protein ser/thr kinase

10 consisting of a kinase domain together with an N-terminal PH domain and C-terminal regulatory domain. The enzyme PKB itself is phosphorylated on Thr 308 by PDK1 and on Ser 473 by an as yet unidentified kinase. Full activation requires phosphorylation at both sites whilst association between PIP3 and the PH domain is required for anchoring of the enzyme to the cytoplasmic face of the lipid membrane

15 providing optimal access to substrates.

Activated PKB in turns phosphorylates a range of substrates contributing to the overall survival response. Whilst we cannot be certain that we understand all of the factors responsible for mediating the PKB dependent survival response, some important actions are believed to be phosphorylation and inactivation of the pro-apoptotic factor BAD and caspase 9, phosphorylation of Forkhead transcription factors e.g. FKHR leading to their exclusion from the nucleus, and activation of the NfkappaB pathway by phosphorylation of upstream kinases in the cascade.

In addition to the anti-apoptotic and pro-survival actions of the PKB pathway, the enzyme also plays an important role in promoting cell proliferation. This action is

25 again likely to be mediated via several actions, some of which are thought to be phosphorylation and inactivation of the cyclin dependent kinase inhibitor of p21<sup>Cip1/WAF1</sup>, and phosphorylation and activation of mTOR, a kinase controlling several aspects of cell growth.

The phosphatase PTEN which dephosphorylates and inactivates polyphosphatidyl-inositols is a key tumour suppressor protein which normally acts to regulate the

PI3K/PKB survival pathway. The significance of the PI3K/PKB pathway in tumourigenesis can be judged from the observation that PTEN is one of the most common targets of mutation in human tumours, with mutations in this phosphatase having been found in ~50% or more of melanomas (Guldberg et al 1997, Cancer Research 57, 3660-3663) and advanced prostate cancers (Cairns et al 1997 Cancer Research 57, 4997). These observations and others suggest that a wide range of tumour types are dependent on the enhanced PKB activity for growth and survival and would respond therapeutically to appropriate inhibitors of PKB.

There are 3 closely related isoforms of PKB called alpha, beta and gamma, which 10 genetic studies suggest have distinct but overlapping functions. Evidence suggests that they can all independently play a role in cancer. For example PKB beta has been found to be over-expressed or activated in 10 – 40% of ovarian and pancreatic cancers (Bellacosa et al 1995, Int. J. Cancer 64, 280 – 285; Cheng et al 1996, PNAS 93, 3636-3641; Yuan et al 2000, Oncogene 19, 2324 – 2330), PKB alpha is 15 amplified in human gastric, prostate and breast cancer (Staal 1987, PNAS 84, 5034 – 5037; Sun et al 2001, Am. J. Pathol. 159, 431 – 437) and increased PKB gamma activity has been observed in steroid independent breast and prostate cell lines (Nakatani et al 1999, J. Biol. Chem. 274, 21528 – 21532).

The PKB pathway also functions in the growth and survival of normal tissues and 20 may be regulated during normal physiology to control cell and tissue function. Thus disorders associated with undesirable proliferation and survival of normal cells and tissues may also benefit therapeutically from treatment with a PKB inhibitor. Examples of such disorders are disorders of immune cells associated with prolonged expansion and survival of cell population leading to a prolonged or up 25 regulated immune response. For example, T and B lymphocyte response to cognate antigens or growth factors such as interferon gamma activates the PI3K/PKB pathway and is responsible for maintaining the survival of the antigen specific lymphocyte clones during the immune response. Under conditions in which lymphocytes and other immune cells are responding to inappropriate self or foreign 30 antigens, or in which other abnormalities lead to prolonged activation, the PKB

pathway contributes an important survival signal preventing the normal mechanisms by which the immune response is terminated via apoptosis of the activated cell population. There is a considerable amount of evidence demonstrating the expansion of lymphocyte populations responding to self antigens 5 in autoimmune conditions such as multiple sclerosis and arthritis. Expansion of lymphocyte populations responding inappropriately to foreign antigens is a feature of another set of conditions such as allergic responses and asthma. In summary inhibition of PKB could provide a beneficial treatment for immune disorders.

Other examples of inappropriate expansion, growth, proliferation, hyperplasia and 10 survival of normal cells in which PKB may play a role include but are not limited to atherosclerosis, cardiac myopathy and glomerulonephritis.

In addition to the role in cell growth and survival, the PKB pathway functions in the control of glucose metabolism by insulin. Available evidence from mice deficient 15 in the alpha and beta isoforms of PKB suggests that this action is mediated by the beta isoform. As a consequence, modulators of PKB activity may also find utility in diseases in which there is a dysfunction of glucose metabolism such as diabetes.

Cyclic AMP-dependent protein kinase (PKA) is a serine/threonine protein kinase that phosphorylates a wide range of substrates and is involved in the regulation of many cellular processes including cell growth, cell differentiation, ion-channel 20 conductivity, gene transcription and synaptic release of neurotransmitters. In its inactive form, the PKA holoenzyme is a tetramer comprising two regulatory subunits and two catalytic subunits.

PKA acts as a link between G-protein mediated signal transduction events and the cellular processes that they regulate. Binding of a hormone ligand such as glucagon 25 to a transmembrane receptor activates a receptor-coupled G-protein (GTP-binding and hydrolyzing protein). Upon activation, the alpha subunit of the G protein dissociates and binds to and activates adenylate cyclase, which in turn converts ATP to cyclic-AMP (cAMP). The cAMP thus produced then binds to the regulatory subunits of PKA leading to dissociation of the associated catalytic subunits. The

catalytic subunits of PKA, which are inactive when associated with the regulatory sub-units, become active upon dissociation and take part in the phosphorylation of other regulatory proteins.

For example, the catalytic sub-unit of PKA phosphorylates the kinase

5 Phosphorylase Kinase which is involved in the phosphorylation of Phosphorylase, the enzyme responsible for breaking down glycogen to release glucose. PKA is also involved in the regulation of glucose levels by phosphorylating and deactivating glycogen synthase. Thus, PKA may be useful in the treatment or management of diseases in which there is a dysfunction of glucose metabolism such

10 as diabetes.

PKA has also been established as an acute inhibitor of T cell activation. Anndahl *et al.* have investigated the possible role of PKA type I in HIV-induced T cell dysfunction on the basis that T cells from HIV-infected patients have increased levels of cAMP and are more sensitive to inhibition by cAMP analogues than are

15 normal T cells. From their studies, they concluded that increased activation of PKA type I may contribute to progressive T cell dysfunction in HIV infection and that PKA type I may therefore be a potential target for immunomodulating therapy.- Aandahl, E. M., Aukrust, P., Skålhegg, B. S., Müller, F., Frøland, S. S., Hansson, V., Taskén, K. *Protein kinase A type I antagonist restores immune responses of T*

20 *cells from HIV-infected patients. FASEB J.* 12, 855-862 (1998).

Because of the diversity and importance of PKA as a messenger in cell regulation, abnormal responses of cAMP leads to a variety of human diseases derived from this, such as irregular cell growth and proliferation (Stratakis, C.A.; Cho-Chung, Y.S.; Protein Kinase A and human diseases. *Trends Endocrinol. Metab.* 2002, 13, 50-52). Over-expression of PKA has been observed in a variety of human cancer cells including those from ovarian, breast and colon patients. Inhibition of PKA would therefore be an approach to treatment of cancer (Li, Q.; Zhu, G-D.; *Current Topics in Medicinal Chemistry*, 2002, 2, 939-971).

For a review of the role of PKA in human disease, see for example, *Protein Kinase A and Human Disease*, Edited by Constantine A. Stratakis, Annals of the New York Academy of Sciences, Volume 968, 2002, ISBN 1-57331-412-9.

5 Several classes of compounds have been disclosed as having PKA and PKB inhibitory activity.

For example, a class of isoquinolinyl-sulphonamido-diamines having PKB inhibitory activity is disclosed in WO 01/91754 (Yissum).

#### Summary of the Invention

10 The invention provides compounds that have protein kinase A (PKA) and protein B (PKB) inhibiting or modulating activity, and which it is envisaged will be useful in preventing or treating disease states or conditions mediated by PKA or PKB.

Accordingly, in one aspect, the invention provides novel compounds of the formula (I) as defined herein.

15 The invention also provides a compound of the formula (I) as defined herein for use in the prophylaxis or treatment of a disease state or condition mediated by protein kinase A or protein kinase B.

The invention also provides the use of a compound of the formula (I) as defined herein for the manufacture of a medicament for the prophylaxis or treatment of a disease state or condition mediated by protein kinase A or protein kinase B.

20 In a further aspect, the invention provides a method for the prophylaxis or treatment of a disease state or condition mediated by protein kinase A or protein kinase B, which method comprises administering to a subject in need thereof a compound of the formula (I) as defined herein.

25 The invention further provides a method for treating a disease or condition comprising or arising from abnormal cell growth in a mammal, the method comprising administering to the mammal a compound of the formula (I) as defined

herein in an amount effective to inhibit protein kinase A or protein kinase B activity.

In another aspect, the invention provides a method of inhibiting protein kinase A or protein kinase B, which method comprises contacting the kinase with a kinase-inhibiting compound of the formula (I) as defined herein.

The invention further provides a method of modulating a cellular process (for example cell division) by inhibiting the activity of a protein kinase A or a protein kinase B using a compound of the formula (I) as defined herein.

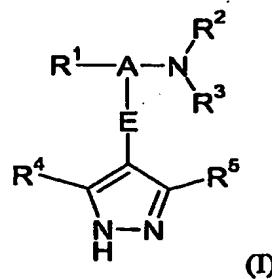
The invention also provides the use of a compound of the formula (I) as defined herein for the manufacture of a medicament for the prophylaxis or treatment of a disease state or condition arising from abnormal cell growth.

The invention also provides a method for treating a disease or condition comprising or arising from abnormal cell growth in a mammal, which method comprises administering to the mammal a compound of the formula (I) as defined herein in an amount effective in inhibiting abnormal cell growth.

In a further aspect, the invention provides a pharmaceutical composition comprising a novel compound of the formula (I) as hereinbefore defined and a pharmaceutically acceptable carrier.

The invention also provides compounds of the formula (I) for use in medicine.

20 The compounds of the invention are represented by the general formula (I):



wherein A is a saturated hydrocarbon linker group containing from 1 to 7 carbon atoms, the linker group having a maximum chain length of 5 atoms extending between R<sup>1</sup> and NR<sup>2</sup>R<sup>3</sup> and a maximum chain length of 4 atoms extending between E and NR<sup>2</sup>R<sup>3</sup>, wherein one of the carbon atoms in the linker group may optionally be replaced by an oxygen or nitrogen atom; and wherein the carbon atoms of the linker group A may optionally bear one or more substituents selected from fluorine and hydroxy, provided that the hydroxy group is not located at a carbon atom  $\alpha$  with respect to the NR<sup>2</sup>R<sup>3</sup> group;

E is a monocyclic or bicyclic carbocyclic or heterocyclic group;

R<sup>1</sup> is an aryl or heteroaryl group;

R<sup>2</sup> and R<sup>3</sup> are independently selected from hydrogen, C<sub>1-4</sub> hydrocarbyl and C<sub>1-4</sub> acyl;

or R<sup>2</sup> and R<sup>3</sup> together with the nitrogen atom to which they are attached form a saturated monocyclic heterocyclic group having 4-7 ring members and optionally containing a second heteroatom ring member selected from O and N;

or one of R<sup>2</sup> and R<sup>3</sup> together with the nitrogen atom to which they are attached and one or more atoms from the linker group A form a saturated monocyclic heterocyclic group having 4-7 ring members and optionally containing a second heteroatom ring member selected from O and N;

or NR<sup>2</sup>R<sup>3</sup> and the carbon atom of linker group A to which it is attached together form a cyano group;

R<sup>4</sup> is selected from hydrogen, halogen, C<sub>1-5</sub> saturated hydrocarbyl, cyano, and CF<sub>3</sub>; and

R<sup>5</sup> is selected from selected from hydrogen, halogen, C<sub>1-5</sub> saturated hydrocarbyl, cyano, CONH<sub>2</sub>, CF<sub>3</sub>, NH<sub>2</sub>, NHCOR<sup>9</sup> or NHCONHR<sup>9</sup>;

R<sup>9</sup> is phenyl or benzyl each optionally substituted by one or substituents selected from halogen, hydroxy, trifluoromethyl, cyano, nitro, carboxy, amino, mono- or di-C<sub>1-4</sub> hydrocarbylamino; a group R<sup>a</sup>-R<sup>b</sup> wherein R<sup>a</sup> is a bond, O, CO, X<sup>1</sup>C(X<sup>2</sup>), C(X<sup>2</sup>)X<sup>1</sup>, X<sup>1</sup>C(X<sup>2</sup>)X<sup>1</sup>, S, SO, SO<sub>2</sub>, NR<sup>c</sup>, SO<sub>2</sub>NR<sup>c</sup> or NR<sup>c</sup>SO<sub>2</sub>; and R<sup>b</sup> is selected from hydrogen, heterocyclic groups having from 3 to 12 ring members, and a C<sub>1-8</sub> hydrocarbyl group optionally substituted by one or more substituents selected

from hydroxy, oxo, halogen, cyano, nitro, carboxy, amino, mono- or di- $C_{1-4}$  hydrocarbylamino, carbocyclic and heterocyclic groups having from 3 to 12 ring members and wherein one or more carbon atoms of the  $C_{1-8}$  hydrocarbyl group may optionally be replaced by O, S, SO, SO<sub>2</sub>, NR<sup>c</sup>, X<sup>1</sup>C(X<sup>2</sup>), C(X<sup>2</sup>)X<sup>1</sup> or X<sup>1</sup>C(X<sup>2</sup>)X<sup>1</sup>;

5        R<sup>c</sup> is selected from hydrogen and  $C_{1-4}$  hydrocarbyl; and  
       X<sup>1</sup> is O, S or NR<sup>c</sup> and X<sup>2</sup> is =O, =S or =NR<sup>c</sup>.

General Preferences and Definitions

The following general preferences and definitions shall apply to each of the  
       10      moieties A, E and R<sup>1</sup> to R<sup>5</sup> and R<sup>9</sup> and any sub-definition, sub-group or embodiment thereof, unless the context indicates otherwise.

References to "carbocyclic" and "heterocyclic" groups as used herein shall, unless the context indicates otherwise, include both aromatic and non-aromatic ring systems. In general, such groups may be monocyclic or bicyclic and may contain, for example, 3 to 12 ring members, more usually 5 to 10 ring members. Examples  
       15      of monocyclic groups are groups containing 3, 4, 5, 6, 7, and 8 ring members, more usually 3 to 7, and preferably 5 or 6 ring members. Examples of bicyclic groups are those containing 8, 9, 10, 11 and 12 ring members, and more usually 9 or 10 ring members.

The carbocyclic or heterocyclic groups can be aryl or heteroaryl groups having  
       20      from 5 to 12 ring members, more usually from 5 to 10 ring members. The term "aryl" as used herein refers to a carbocyclic group having aromatic character and the term "heteroaryl" is used herein to denote a heterocyclic group having aromatic character. The terms "aryl" and "heteroaryl" embrace polycyclic (e.g. bicyclic) ring systems wherein one or more rings are non-aromatic, provided that at least one ring  
       25      is aromatic. In such polycyclic systems, the group may be attached by the aromatic ring, or by a non-aromatic ring. The aryl or heteroaryl groups can be monocyclic or bicyclic groups and can be unsubstituted or substituted with one or more substituents, for example one or more groups R<sup>10</sup> as defined herein.

The term non-aromatic group embraces unsaturated ring systems without aromatic character, partially saturated and fully saturated carbocyclic and heterocyclic ring systems. The terms "unsaturated" and "partially saturated" refer to rings wherein the ring structure(s) contains atoms sharing more than one valence bond i.e. the ring

5 contains at least one multiple bond e.g. a C=C, C≡C or N=C bond. The term "fully saturated" refers to rings where there are no multiple bonds between ring atoms. Saturated carbocyclic groups include cycloalkyl groups as defined below. Partially saturated carbocyclic groups include cycloalkenyl groups as defined below, for example cyclopentenyl, cycloheptenyl and cyclooctenyl.

10 Examples of heteroaryl groups are monocyclic and bicyclic groups containing from five to twelve ring members, and more usually from five to ten ring members. The heteroaryl group can be, for example, a five membered or six membered monocyclic ring or a bicyclic structure formed from fused five and six membered rings or two fused six membered rings. Each ring may contain up to about four

15 heteroatoms typically selected from nitrogen, sulphur and oxygen. Typically the heteroaryl ring will contain up to 3 heteroatoms, more usually up to 2, for example a single heteroatom. In one embodiment, the heteroaryl ring contains at least one ring nitrogen atom. The nitrogen atoms in the heteroaryl rings can be basic, as in the case of an imidazole or pyridine, or essentially non-basic as in the case of an

20 indole or pyrrole nitrogen. In general the number of basic nitrogen atoms present in the heteroaryl group, including any amino group substituents of the ring, will be less than five.

Examples of heteroaryl groups include but are not limited to pyridine, pyrrole, furan, thiophene, imidazole, furazan, oxazole, oxadiazole, oxatriazole, isoxazole, thiazole, isothiazole, pyrazole, pyrazine, pyridazine, pyrimidine, triazine, triazole, 25 tetrazole, quinoline, isoquinoline, benzfuran, benzthiophene, chroman, thiochroman, benzimidazole, benzoxazole, benzisoxazole, benzthiazole, benzisothiazole, isobenzofuran, indole, isoindole, indolizine, indoline, isoindoline, purine (e.g., adenine, guanine), indazole, benzodioxole, chromene, isochromene, 30 chroman, isochroman, benzodioxan, quinolizine, benzoxazine, benzodiazine,

pyridopyridine, pyrazolopyridine, quinoxaline, quinazoline, cinnoline, phthalazine, naphthyridine and pteridine groups.

Examples of polycyclic aryl and heteroaryl groups containing an aromatic ring and a non-aromatic ring include tetrahydronaphthalene, tetrahydroisoquinoline,

5 tetrahydroquinoline, dihydrobenzthiene, dihydrobenzfuran, 2,3-dihydro-benzo[1,4]dioxine, benzo[1,3]dioxole, 4,5,6,7-tetrahydrobenzofuran, indoline and indane groups.

Examples of carbocyclic aryl groups include phenyl, naphthyl, indenyl, and tetrahydronaphthyl groups.

10 Examples of non-aromatic heterocyclic groups are groups having from 3 to 12 ring members, more usually 5 to 10 ring members. Such groups can be monocyclic or bicyclic, for example, and typically have from 1 to 5 heteroatom ring members (more usually 1, 2, 3 or 4 heteroatom ring members), usually selected from nitrogen, oxygen and sulphur. The heterocyclic groups can contain, for example,

15 cyclic ether moieties (e.g. as in tetrahydrofuran and dioxane), cyclic thioether moieties (e.g. as in tetrahydrothiophene and dithiane), cyclic amine moieties (e.g. as in pyrrolidine), cyclic sulphones (e.g. as in sulfolane and sulfolene), cyclic sulphoxides, cyclic sulphonamides and combinations thereof (e.g. thiomorpholine).

Particular examples include morpholine, piperidine (e.g. 1-piperidinyl, 2-piperidinyl 3-piperidinyl and 4-piperidinyl), piperidone, pyrrolidine (e.g. 1-pyrrolidinyl, 2-pyrrolidinyl and 3-pyrrolidinyl), pyrrolidone, pyran (2H-pyran or 4H-pyran), dihydrothiophene, dihydropyran, dihydrofuran, dihydrothiazole, tetrahydrofuran, tetrahydrothiophene, dioxane, tetrahydropyran (e.g. 4-tetrahydro pyranyl), imidazoline, imidazolidinone, oxazoline, thiazoline, 2-pyrazoline, 25 pyrazolidine, piperazone, piperazine, and N-alkyl piperazines such as N-methyl piperazine. In general, preferred non-aromatic heterocyclic groups include piperidine, pyrrolidine, morpholine, piperazine and N-alkyl piperazines.

Examples of non-aromatic carbocyclic groups include cycloalkane groups such as cyclohexyl and cyclopentyl, cycloalkenyl groups such as cyclopentenyl, cyclohexenyl, cycloheptenyl and cyclooctenyl, as well as cyclohexadienyl, cyclooctatetraene, tetrahydronaphthyl and decalinyl.

5 Where reference is made herein to carbocyclic and heterocyclic groups, the carbocyclic or heterocyclic ring can, unless the context indicates otherwise, be unsubstituted or substituted by one or more substituent groups  $R^{10}$  selected from halogen, hydroxy, trifluoromethyl, cyano, nitro, carboxy, amino, mono- or di- $C_{1-4}$  hydrocarbyl amino, carbocyclic and heterocyclic groups having from 3 to 12 ring members; a group  $R^a$ - $R^b$  wherein  $R^a$  is a bond, O, CO,  $X^1C(X^2)$ ,  $C(X^2)X^1$ ,  $X^1C(X^2)X^1$ , S, SO,  $SO_2$ ,  $NR^c$ ,  $SO_2NR^c$  or  $NR^cSO_2$ ; and  $R^b$  is selected from hydrogen, carbocyclic and heterocyclic groups having from 3 to 12 ring members, and a  $C_{1-8}$  hydrocarbyl group optionally substituted by one or more substituents selected from hydroxy, oxo, halogen, cyano, nitro, carboxy, amino, mono- or di- $C_{1-4}$  hydrocarbyl amino, carbocyclic and heterocyclic groups having from 3 to 12 ring members and wherein one or more carbon atoms of the  $C_{1-8}$  hydrocarbyl group may optionally be replaced by O, S, SO,  $SO_2$ ,  $NR^c$ ,  $X^1C(X^2)$ ,  $C(X^2)X^1$  or  $X^1C(X^2)X^1$ ;

10  $R^c$  is selected from hydrogen and  $C_{1-4}$  hydrocarbyl; and

15  $X^1$  is O, S or  $NR^c$  and  $X^2$  is =O, =S or = $NR^c$ .

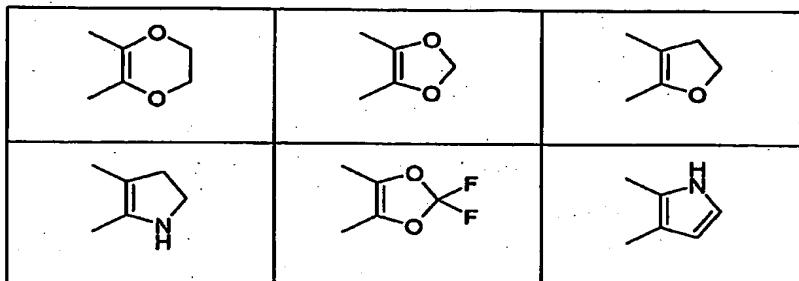
20 Where the substituent group  $R^{10}$  comprises or includes a carbocyclic or heterocyclic group, the said carbocyclic or heterocyclic group may be unsubstituted or may itself be substituted with one or more further substituent groups  $R^{10}$ . In one sub-group of compounds of the formula (I), such further substituent groups  $R^{10}$  may include

25 carbocyclic or heterocyclic groups, which are typically not themselves further substituted. In another sub-group of compounds of the formula (I), the said further substituents do not include carbocyclic or heterocyclic groups but are otherwise selected from the groups listed above in the definition of  $R^{10}$ .

30 Where the carbocyclic and heterocyclic groups have a pair of substituents on adjacent ring atoms, the two substituents may be linked so as to form a cyclic

group. For example, an adjacent pair of substituents on adjacent carbon atoms of a ring may be linked via one or more heteroatoms and optionally substituted alkylene groups to form a fused oxa-, dioxa-, aza-, diaza- or oxa-aza-cycloalkyl group.

Examples of such linked substituent groups include:



5 Examples of halogen substituents include fluorine, chlorine, bromine and iodine. Fluorine and chlorine are particularly preferred.

In the definition of the compounds of the formula (I) above and as used hereinafter, the term "hydrocarbyl" is a generic term encompassing aliphatic, alicyclic and aromatic groups having an all-carbon backbone, except where otherwise stated. In

10 certain cases, as defined herein, one or more of the carbon atoms making up the carbon backbone may be replaced by a specified atom or group of atoms. Examples of hydrocarbyl groups include alkyl, cycloalkyl, cycloalkenyl, carbocyclic aryl, alkenyl, alkynyl, cycloalkylalkyl, cycloalkenylalkyl, and carbocyclic aralkyl, aralkenyl and aralkynyl groups. Such groups can be

15 unsubstituted or, where stated, can be substituted by one or more substituents as defined herein. The examples and preferences expressed below apply to each of the hydrocarbyl substituent groups or hydrocarbyl-containing substituent groups referred to in the various definitions of substituents for compounds of the formula (I) unless the context indicates otherwise.

20 Generally by way of example, the hydrocarbyl groups can have up to eight carbon atoms, unless the context requires otherwise. Within the sub-set of hydrocarbyl groups having 1 to 8 carbon atoms, particular examples are C<sub>1-6</sub> hydrocarbyl groups, such as C<sub>1-4</sub> hydrocarbyl groups (e.g. C<sub>1-3</sub> hydrocarbyl groups or C<sub>1-2</sub>

hydrocarbyl groups), specific examples being any individual value or combination of values selected from C<sub>1</sub>, C<sub>2</sub>, C<sub>3</sub>, C<sub>4</sub>, C<sub>5</sub>, C<sub>6</sub>, C<sub>7</sub> and C<sub>8</sub> hydrocarbyl groups.

The term "alkyl" covers both straight chain and branched chain alkyl groups.

Examples of alkyl groups include methyl, ethyl, propyl, isopropyl, n-butyl, isobutyl, 5 tert-butyl, n-pentyl, 2-pentyl, 3-pentyl, 2-methyl butyl, 3-methyl butyl, and n-hexyl and its isomers. Within the sub-set of alkyl groups having 1 to 8 carbon atoms, particular examples are C<sub>1-6</sub> alkyl groups, such as C<sub>1-4</sub> alkyl groups (e.g. C<sub>1-3</sub> alkyl groups or C<sub>1-2</sub> alkyl groups).

10 Examples of cycloalkyl groups are those derived from cyclopropane, cyclobutane, cyclopentane, cyclohexane and cycloheptane. Within the sub-set of cycloalkyl groups the cycloalkyl group will have from 3 to 8 carbon atoms, particular examples being C<sub>3-6</sub> cycloalkyl groups.

15 Examples of alkenyl groups include, but are not limited to, ethenyl (vinyl), 1-propenyl, 2-propenyl (allyl), isopropenyl, butenyl, buta-1,4-dienyl, pentenyl, and hexenyl. Within the sub-set of alkenyl groups the alkenyl group will have 2 to 8 carbon atoms, particular examples being C<sub>2-6</sub> alkenyl groups, such as C<sub>2-4</sub> alkenyl groups.

20 Examples of cycloalkenyl groups include, but are not limited to, cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclopentadienyl and cyclohexenyl. Within the sub-set of cycloalkenyl groups the cycloalkenyl groups have from 3 to 8 carbon atoms, and particular examples are C<sub>3-6</sub> cycloalkenyl groups.

Examples of alkynyl groups include, but are not limited to, ethynyl and 2-propynyl (propargyl) groups. Within the sub-set of alkynyl groups having 2 to 8 carbon atoms, particular examples are C<sub>2-6</sub> alkynyl groups, such as C<sub>2-4</sub> alkynyl groups.

25 Examples of carbocyclic aryl groups include substituted and unsubstituted phenyl, naphthyl, indane and indene groups.

Examples of cycloalkylalkyl, cycloalkenylalkyl, carbocyclic aralkyl, aralkenyl and aralkynyl groups include phenethyl, benzyl, styryl, phenylethynyl, cyclohexylmethyl, cyclopentylmethyl, cyclobutylmethyl, cyclopropylmethyl and cyclopentenylmethyl groups.

5 When present, a hydrocarbyl group can be optionally substituted by one or more substituents selected from hydroxy, oxo, alkoxy, carboxy, halogen, cyano, nitro, amino, mono- or di-C<sub>1-4</sub> hydrocarbylamino, and monocyclic or bicyclic carbocyclic and heterocyclic groups having from 3 to 12 (typically 3 to 10 and more usually 5 to 10) ring members. Preferred substituents include halogen such as fluorine.

10 Thus, for example, the substituted hydrocarbyl group can be a partially fluorinated or perfluorinated group such as difluoromethyl or trifluoromethyl. In one embodiment preferred substituents include monocyclic carbocyclic and heterocyclic groups having 3-7 ring members.

One or more carbon atoms of a hydrocarbyl group may optionally be replaced by O, 15 S, SO, SO<sub>2</sub>, NR<sup>c</sup>, X<sup>1</sup>C(X<sup>2</sup>), C(X<sup>2</sup>)X<sup>1</sup> or X<sup>1</sup>C(X<sup>2</sup>)X<sup>1</sup> wherein X<sup>1</sup> and X<sup>2</sup> are as hereinbefore defined. For example, 1, 2, 3 or 4 carbon atoms of the hydrocarbyl group may be replaced by one of the atoms or groups listed, and the replacing atoms or groups may be the same or different. Examples of groups in which a carbon atom of the hydrocarbyl group has been replaced by a replacement atom or 20 group as defined above include ethers and thioethers (C replaced by O or S), amides, esters, thioamides and thioesters (C replaced by X<sup>1</sup>C(X<sup>2</sup>) or C(X<sup>2</sup>)X<sup>1</sup>), sulphones and sulfoxides (C replaced by SO or SO<sub>2</sub>) and amines (C replaced by NR<sup>c</sup>).

Where an amino group has two hydrocarbyl substituents, they may, together with 25 the nitrogen atom to which they are attached, and optionally with another heteroatom such as nitrogen, sulphur, or oxygen, link to form a ring structure of 4 to 7 ring members.

The definition "R<sup>a</sup>-R<sup>b</sup>" as used herein, either with regard to substituents present on a carbocyclic or heterocyclic moiety, or with regard to other substituents present at

other locations on the compounds of the formula (I), includes *inter alia* compounds wherein R<sup>a</sup> is selected from a bond, O, CO, OC(O), SC(O), NR<sup>c</sup>C(O), OC(S), SC(S), NR<sup>c</sup>C(S), OC(NR<sup>c</sup>), SC(NR<sup>c</sup>), NR<sup>c</sup>C(NR<sup>c</sup>), C(O)O, C(O)S, C(O)NR<sup>c</sup>, C(S)O, C(S)S, C(S) NR<sup>c</sup>, C(NR<sup>c</sup>)O, C(NR<sup>c</sup>)S, C(NR<sup>c</sup>)NR<sup>c</sup>, OC(O)O, SC(O)O, 5 NR<sup>c</sup>C(O)O, OC(S)O, SC(S)O, NR<sup>c</sup>C(S)O, OC(NR<sup>c</sup>)O, SC(NR<sup>c</sup>)O, NR<sup>c</sup>C(NR<sup>c</sup>)O, OC(O)S, SC(O)S, NR<sup>c</sup>C(O)S, OC(S)S, SC(S)S, NR<sup>c</sup>C(S)S, OC(NR<sup>c</sup>)S, SC(NR<sup>c</sup>)S, NR<sup>c</sup>C(NR<sup>c</sup>)S, OC(O)NR<sup>c</sup>, SC(O)NR<sup>c</sup>, NR<sup>c</sup>C(O) NR<sup>c</sup>, OC(S)NR<sup>c</sup>, SC(S) NR<sup>c</sup>, NR<sup>c</sup>C(S)NR<sup>c</sup>, OC(NR<sup>c</sup>)NR<sup>c</sup>, SC(NR<sup>c</sup>)NR<sup>c</sup>, NR<sup>c</sup>C(NR<sup>c</sup>)NR<sup>c</sup>, S, SO, SO<sub>2</sub>, NR<sup>c</sup>, SO<sub>2</sub>NR<sup>c</sup> and NR<sup>c</sup>SO<sub>2</sub> wherein R<sup>c</sup> is as hereinbefore defined.

10 The moiety R<sup>b</sup> can be hydrogen or it can be a group selected from carbocyclic and heterocyclic groups having from 3 to 12 ring members (typically 3 to 10 and more usually from 5 to 10), and a C<sub>1-8</sub> hydrocarbyl group optionally substituted as hereinbefore defined. Examples of hydrocarbyl, carbocyclic and heterocyclic groups are as set out above.

15 Specific Embodiments of and Preferences for A, E, R<sup>1</sup> to R<sup>5</sup> and R<sup>9</sup>

In formula (I), A is a saturated hydrocarbon linker group containing from 1 to 7 carbon atoms, the linker group having a maximum chain length of 5 atoms extending between R<sup>1</sup> and NR<sup>2</sup>R<sup>3</sup> and a maximum chain length of 4 atoms extending between E and NR<sup>2</sup>R<sup>3</sup>. The term "maximum chain length" refers to the 20 number of atoms lying directly between the two moieties in question, and does not take into account any branching in the chain or any hydrogen atoms that may be present. For example, in the structure A shown below:

(A)

the chain length between R<sup>1</sup> and NR<sup>2</sup>R<sup>3</sup> is 3 atoms whereas the chain length 25 between E and NR<sup>2</sup>R<sup>3</sup> is 2 atoms.

In general it is presently preferred that the linker group has a maximum chain length of 3 atoms (more preferably 1 or 2 atoms, and most preferably 2 atoms) extending between  $R^1$  and  $NR^2R^3$ .

It is preferred that the linker group has a maximum chain length of 3 atoms  
5 extending between E and  $NR^2R^3$ .

In one particularly preferred group of compounds, the linker group has a chain length of 2 or 3 atoms extending between  $R^1$  and  $NR^2R^3$  and a chain length of 2 or 3 atoms extending between E and  $NR^2R^3$ .

One of the carbon atoms in the linker group may optionally be replaced by an  
10 oxygen or nitrogen atom, with nitrogen currently being preferred. When present, the nitrogen atom preferably is linked directly to the group E.

When a nitrogen atom or oxygen atom are present, it is preferred that the nitrogen or oxygen atom and the  $NR^2R^3$  group are spaced apart by at least two intervening carbon atoms.

15 In one particular group of compounds within formula (I), the linker atom linked directly to the group E is a carbon atom and the linker group A has an all-carbon skeleton.

The carbon atoms of the linker group A may optionally bear one or more substituents selected from fluorine and hydroxy, provided that the hydroxy group is  
20 not located at a carbon atom  $\alpha$  with respect to the  $NR^2R^3$  group. Typically, the hydroxy group, if present, is located at a position  $\beta$  with respect to the  $NR^2R^3$  group. In general, no more than one hydroxy group will be present. Where fluorine atoms are present, they may be present in a difluoromethylene or trifluoromethyl group, for example.

25 In one embodiment of the invention, no fluorine atoms are present in the linker group A.

In another embodiment of the invention, no hydroxy groups are present in the linker group A.

In one group of compounds of the formula (I) neither hydroxy groups nor fluorine atoms are present in the linker group A, i.e. the linker group A is unsubstituted.

5 In order to modify the susceptibility of the compounds to metabolic degradation *in vivo*, the linker group A can have a branched configuration at the carbon atom attached to the NR<sup>2</sup>R<sup>3</sup> group. For example, the carbon atom attached to the NR<sup>2</sup>R<sup>3</sup> group can be attached to a pair of *gem*-dimethyl groups.

10 In one particular group of compounds of the formula (I), the portion R<sup>1</sup>-A-NR<sup>2</sup>R<sup>3</sup> of the compound is represented by the formula R<sup>1</sup>-(G)<sub>k</sub>-(CH<sub>2</sub>)<sub>m</sub>-X-(CH<sub>2</sub>)<sub>n</sub>-(CR<sup>6</sup>R<sup>7</sup>)<sub>p</sub>-NR<sup>2</sup>R<sup>3</sup> wherein G is NH, NMe or O; X is attached to the group E and is selected from (CH<sub>2</sub>)<sub>j</sub>-CH, (CH<sub>2</sub>)<sub>j</sub>-N and (NH)<sub>j</sub>-CH; , j is 0 or 1, k is 0 or 1, m is 0 or 1, n is 0, 1, 2, or 3 and p is 0 or 1, and the sum of j, k, m, n and p does not exceed 4; and R<sup>6</sup> and R<sup>7</sup> are the same or different and are selected from methyl and ethyl, or CR<sup>6</sup>R<sup>7</sup> forms a cyclopropyl group.

A preferred group CR<sup>6</sup>R<sup>7</sup> is C(CH<sub>3</sub>)<sub>2</sub>.

Preferably X is (CH<sub>2</sub>)<sub>j</sub>-CH.

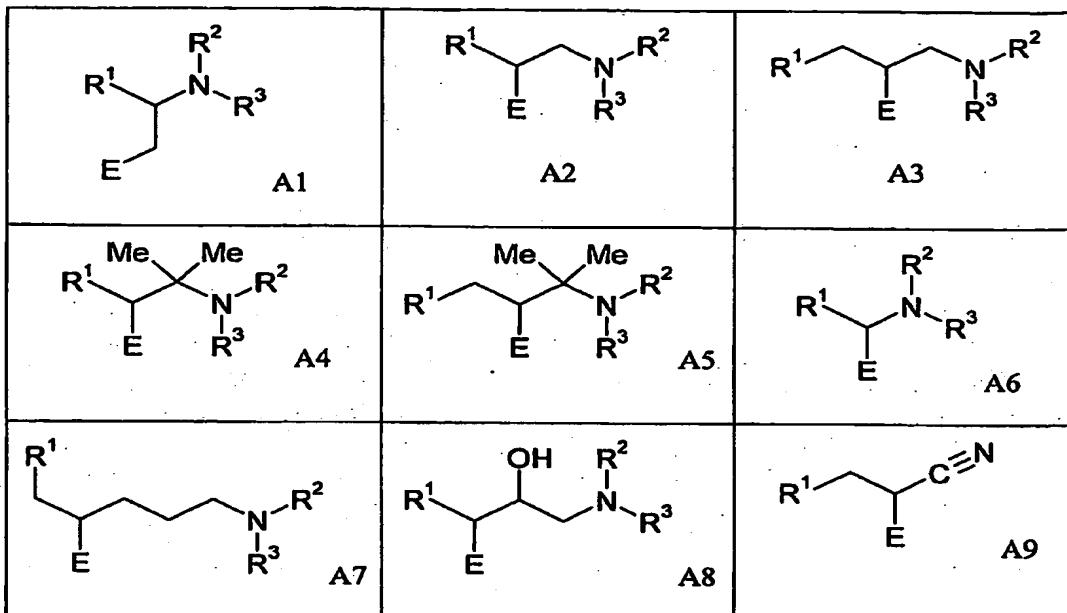
In one preferred configuration, k is 0, m is 0 or 1, n is 0, 1, 2 or 3 and p is 0.

In another preferred configuration, k is 0, m is 0 or 1, n is 0, 1 or 2 and p is 1.

20 In another configuration, X is (CH<sub>2</sub>)<sub>j</sub>-CH, k is 1, m is 0, n is 0, 1,2 or 3 and p is 0.

In another preferred configuration, X is (CH<sub>2</sub>)<sub>j</sub>-CH, k is 1, m is 0, n is 0, 1 or 2 and p is 1.

Particular examples of the linker group A, together with their points of attachment to the groups R<sup>1</sup>, E and NR<sup>2</sup>R<sup>3</sup>, are shown in Table 1 below.



Currently preferred groups include A1, A2 and A3.

In formula (I), E is a monocyclic or bicyclic carbocyclic or heterocyclic group and can be selected from the groups set out above in the section headed General Preferences and Definitions. Preferably E is a monocyclic group.

5 Particular examples of monocyclic groups include monocyclic aryl and heteroaryl groups such as phenyl, thiophene, furan, pyrimidine and pyridine, phenyl being presently preferred.

10 Examples of non-aromatic monocyclic groups include cycloalkanes such as cyclohexane and cyclopentane, and nitrogen-containing rings such as piperazine and piperazone.

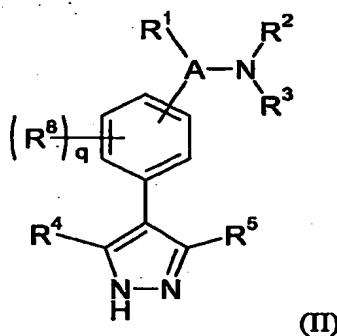
15 It is preferred that the group A and the pyrazole group are attached to the group E in a *meta* or *para* relative orientation; i.e. A and the pyrazole group are not attached to adjacent ring members of the group E. Examples of groups such groups E include 1,4-phenylene, 1,3-phenylene, 2,5-pyridylene and 2,4-pyridylene, 1,4-piperazinyl, and 1,4-piperazonyl.

The groups E can be unsubstituted or can have up to 4 substituents R<sup>8</sup> which may be selected from the group R<sup>10</sup> as hereinbefore defined. More typically however, the substituents R<sup>8</sup> are selected from hydroxy, oxo (when E is non-aromatic), chlorine, bromine, trifluoromethyl, cyano, C<sub>1-4</sub> hydrocarbyloxy and C<sub>1-4</sub> hydrocarbyl

5     optionally substituted by C<sub>1-2</sub> alkoxy or hydroxy.

Preferably there are 0-3 substituents, more preferably 0-2 substituents, for example 0 or 1 substituent. The group E is preferably unsubstituted.

One sub-group of compounds of the formula (I) has the general formula (II):



10     wherein the group A is attached to the *meta* or *para* position of the benzene ring, q is 0-4 and R<sup>8</sup> is a substituent group as hereinbefore defined. In formula (II), q is preferably 0, 1 or 2, more preferably 0 or 1 and most preferably 0.

The group R<sup>1</sup> is an aryl or heteroaryl group and may be selected from the list of such groups set out in the section headed General Preferences and Definitions.

15     Examples of such groups include phenyl, naphthyl, thiienyl, furan, pyrimidine and pyridine, with phenyl being presently preferred.

The group R<sup>1</sup> can be unsubstituted or substituted by up to 5 substituents, and examples of substituents are those listed in group R<sup>10</sup> above. Preferred substituents include hydroxy, C<sub>1-4</sub> acyloxy, fluorine, chlorine, bromine, trifluoromethyl, cyano, C<sub>1-4</sub> hydrocarbyloxy and C<sub>1-4</sub> hydrocarbyl optionally substituted by C<sub>1-2</sub> alkoxy or

hydroxy. Although up to 5 substituents may be present, more typically there are 0, 1, 2, 3 or 4 substituents, preferably 0, 1, 2 or 3, and more preferably 0, 1 or 2.

In one embodiment, the group R<sup>1</sup> can have one or two substituents selected from fluorine, chlorine, trifluoromethyl, methyl and methoxy. When R<sup>1</sup> is a phenyl group, particular examples of substituent combinations include mono-chlorophenyl and dichlorophenyl.

In formula (I), R<sup>4</sup> is selected from hydrogen, halogen, C<sub>1-5</sub> saturated hydrocarbyl, cyano and CF<sub>3</sub>. Preferred values for R<sup>4</sup> include hydrogen and methyl.

In formula (I), R<sup>5</sup> is selected from selected from hydrogen, halogen, C<sub>1-5</sub> saturated hydrocarbyl, cyano, CONH<sub>2</sub>, CF<sub>3</sub>, NH<sub>2</sub>, NHCOR<sup>9</sup> and NHCONHR<sup>9</sup> where R<sup>9</sup> is optionally substituted phenyl or benzyl.

More preferably, R<sup>5</sup> is selected from selected from hydrogen, halogen, C<sub>1-5</sub> saturated hydrocarbyl, cyano, CF<sub>3</sub>, NH<sub>2</sub>, NHCOR<sup>9</sup> and NHCONHR<sup>9</sup> where R<sup>9</sup> is optionally substituted phenyl or benzyl.

15 Particular examples of the moiety R<sup>5</sup> include hydrogen, fluorine, chlorine, bromine, methyl, ethyl, hydroxyethyl, methoxymethyl, cyano, CF<sub>3</sub>, NH<sub>2</sub>, NHCOR<sup>9a</sup> and NHCONHR<sup>9a</sup> where R<sup>9a</sup> is phenyl or benzyl optionally substituted by hydroxy, C<sub>1-4</sub> acyloxy, fluorine, chlorine, bromine, trifluoromethyl, cyano, C<sub>1-4</sub> hydrocarbyloxy and C<sub>1-4</sub> hydrocarbyl optionally substituted by C<sub>1-2</sub> alkoxy or hydroxy.

20 For the avoidance of doubt, it is to be understood that each general and specific preference, embodiment and example of the groups R<sup>1</sup> may be combined with each general and specific preference, embodiment and example of the groups R<sup>2</sup> and/or R<sup>3</sup> and/or R<sup>4</sup> and/or R<sup>5</sup> and/or R<sup>9</sup> and that all such combinations are embraced by this application.

25 The various functional groups and substituents making up the compounds of the formula (I) are typically chosen such that the molecular weight of the compound of the formula (I) does not exceed 1000. More usually, the molecular weight of the

compound will be less than 750, for example less than 700, or less than 650, or less than 600, or less than 550. More preferably, the molecular weight is less than 525 and, for example, is 500 or less.

Particular compounds of the invention are as illustrated in the examples below.

- 5 Many compounds of the formula (I) can exist in the form of salts, for example acid addition salts or, in certain cases salts of organic and inorganic bases such as carboxylate, sulphonate and phosphate salts. All such salts are within the scope of this invention, and references to compounds of the formula (I) include the salt forms of the compounds.
- 10 Acid addition salts may be formed with a wide variety of acids, both inorganic and organic. Examples of acid addition salts include salts formed with hydrochloric, hydriodic, phosphoric, nitric, sulphuric, citric, lactic, succinic, maleic, malic, isethionic, fumaric, benzenesulphonic, toluenesulphonic, methanesulphonic, ethanesulphonic, naphthalenesulphonic, valeric, acetic, propanoic, butanoic, 15 malonic, glucuronic and lactobionic acids.

If the compound is anionic, or has a functional group which may be anionic (e.g., -COOH may be -COO<sup>-</sup>), then a salt may be formed with a suitable cation.

Examples of suitable inorganic cations include, but are not limited to, alkali metal ions such as Na<sup>+</sup> and K<sup>+</sup>, alkaline earth cations such as Ca<sup>2+</sup> and Mg<sup>2+</sup>, and other cations such as Al<sup>3+</sup>. Examples of suitable organic cations include, but are not limited to, ammonium ion (i.e., NH<sub>4</sub><sup>+</sup>) and substituted ammonium ions (e.g., NH<sub>3</sub>R<sup>+</sup>, NH<sub>2</sub>R<sub>2</sub><sup>+</sup>, NHR<sub>3</sub><sup>+</sup>, NR<sub>4</sub><sup>+</sup>). Examples of some suitable substituted ammonium ions are those derived from: ethylamine, diethylamine, dicyclohexylamine, triethylamine, butylamine, ethylenediamine, ethanolamine, diethanolamine, 20 piperazine, benzylamine, phenylbenzylamine, choline, meglumine, and tromethamine, as well as amino acids, such as lysine and arginine. An example of a common quaternary ammonium ion is N(CH<sub>3</sub>)<sub>4</sub><sup>+</sup>.

Where the compounds of the formula (I) contain an amine function, these may form quaternary ammonium salts, for example by reaction with an alkylating agent according to methods well known to the skilled person. Such quaternary ammonium compounds are within the scope of formula (I).

5. Compounds of the formula (I) containing an amine function may also form N-oxides. A reference herein to a compound of the formula (I) that contains an amine function also includes the N-oxide.

Where a compound contains several amine functions, one or more than one nitrogen atom may be oxidised to form an N-oxide. Particular examples of N-oxides are the N-oxides of a tertiary amine or a nitrogen atom of a nitrogen-containing heterocycle.

- 10

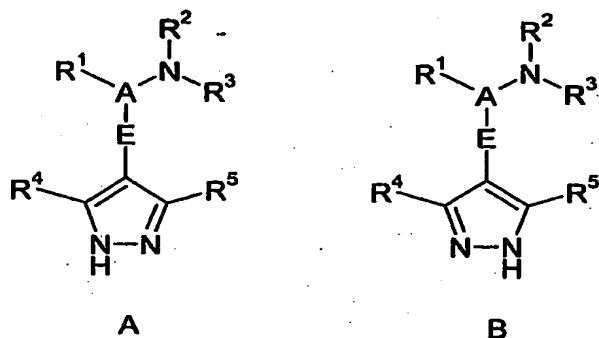
N-Oxides can be formed by treatment of the corresponding amine with an oxidizing agent such as hydrogen peroxide or a per-acid (e.g. a peroxy carboxylic acid), see for example *Advanced Organic Chemistry*, by Jerry March, 4<sup>th</sup> Edition, Wiley 15 Interscience, pages. More particularly, N-oxides can be made by the procedure of L. W. Deady (*Syn. Comm.* 1977, 7, 509-514) in which the amine compound is reacted with *m*-chloroperoxybenzoic acid (MCPBA), for example, in an inert solvent such as dichloromethane.

- 20

Compounds of the formula may exist in a number of different geometric isomeric, and tautomeric forms and references to compounds of the formula (I) include all such forms. For the avoidance of doubt, where a compound can exist in one of several geometric isomeric or tautomeric forms and only one is specifically described or shown, all others are nevertheless embraced by formula (I).

- 25

For example, in compounds of the formula (I) the pyrazole group may take either of the following two tautomeric forms A and B.



For simplicity, the general formula (I) illustrates form A but the formula is to be taken as embracing both form A and form B.

5      Esters such as carboxylic acid esters and acyloxy esters of the compounds of formula (I) bearing a carboxylic acid group or a hydroxyl group are also embraced by Formula (I). Examples of esters are compounds containing the group -C(=O)OR, wherein R is an ester substituent, for example, a C<sub>1-7</sub> alkyl group, a C<sub>3-20</sub> heterocyclyl group, or a C<sub>5-20</sub> aryl group, preferably a C<sub>1-7</sub> alkyl group. Particular examples of ester groups include, but are not limited to, -C(=O)OCH<sub>3</sub>,

10     -C(=O)OCH<sub>2</sub>CH<sub>3</sub>, -C(=O)OC(CH<sub>3</sub>)<sub>3</sub>, and -C(=O)OPh. Examples of acyloxy (reverse ester) groups are represented by -OC(=O)R, wherein R is an acyloxy substituent, for example, a C<sub>1-7</sub> alkyl group, a C<sub>3-20</sub> heterocyclyl group, or a C<sub>5-20</sub> aryl group, preferably a C<sub>1-7</sub> alkyl group. Particular examples of acyloxy groups include, but are not limited to, -OC(=O)CH<sub>3</sub> (acetoxy), -OC(=O)CH<sub>2</sub>CH<sub>3</sub>,

15     -OC(=O)C(CH<sub>3</sub>)<sub>3</sub>, -OC(=O)Ph, and -OC(=O)CH<sub>2</sub>Ph.

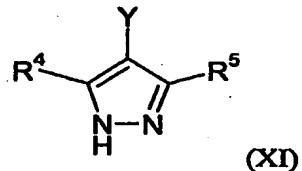
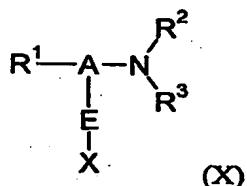
Also encompassed by formula (I) are any polymorphic forms of the compounds, solvates (e.g. hydrates), complexes (e.g. inclusion complexes or clathrates with compounds such as cyclodextrins, or complexes with metals) of the compounds, and pro-drugs of the compounds. By "prodrugs" is meant for example any compound that is converted *in vivo* into a biologically active compound of the formula (I).

For example, some prodrugs are esters of the active compound (e.g., a physiologically acceptable metabolically labile ester). During metabolism, the ester group (-C(=O)OR) is cleaved to yield the active drug. Such esters may be formed by esterification, for example, of any of the carboxylic acid groups (-C(=O)OH) in the parent compound, with, where appropriate, prior protection of any other reactive groups present in the parent compound, followed by deprotection if required.

5 Examples of such metabolically labile esters include those of the formula -  
C(=O)OR wherein R is:  
C<sub>1-7</sub>alkyl (e.g., -Me, -Et, -nPr, -iPr, -nBu, -sBu, -iBu, -tBu);  
10 C<sub>1-7</sub>aminoalkyl (e.g., aminoethyl; 2-(N,N-diethylamino)ethyl;  
2-(4-morpholino)ethyl); and  
acyloxy-C<sub>1-7</sub>alkyl (e.g., acyloxymethyl; acyloxyethyl; pivaloyloxymethyl;  
acetoxyethyl; 1-acetoxyethyl; 1-(1-methoxy-1-methyl)ethyl-carbonyloxyethyl; 1-(benzoyloxy)ethyl; isopropoxy-carbonyloxyethyl; 1-isopropoxy-  
15 carbonyloxyethyl; cyclohexyl-carbonyloxyethyl; 1-cyclohexyl-carbonyloxyethyl;  
cyclohexyloxy-carbonyloxyethyl; 1-cyclohexyloxy-carbonyloxyethyl; (4-tetrahydropyranyloxy) carbonyloxymethyl; 1-(4-tetrahydropyranyloxy)-  
carbonyloxyethyl; (4-tetrahydropyranyl)carbonyloxymethyl; and  
1-(4-tetrahydropyranyl)-carbonyloxyethyl).  
20 Also, some prodrugs are activated enzymatically to yield the active compound, or a compound which, upon further chemical reaction, yields the active compound (for example, as in antigen-directed enzyme pro-drug therapy (ADEPT), gene-directed enzyme pro-drug therapy (GDEPT) and ligand-directed enzyme pro-drug therapy (LIDEP). For example, the prodrug may be a sugar derivative or other glycoside  
25 conjugate, or may be an amino acid ester derivative.

#### Methods for the preparation of compounds of the formula (I)

Compounds of the formula (I) can be prepared by reaction of a compound of the formula (X) with a compound of the formula (XI) or an N-protected derivative thereof:



wherein A, E, and R<sup>1</sup> to R<sup>5</sup> are as hereinbefore defined, one of the groups X and Y is chlorine, bromine or iodine or a trifluoromethanesulphonate (triflate) group, and the other one of the groups X and Y is a boronate residue, for example a boronate ester or boronic acid residue.

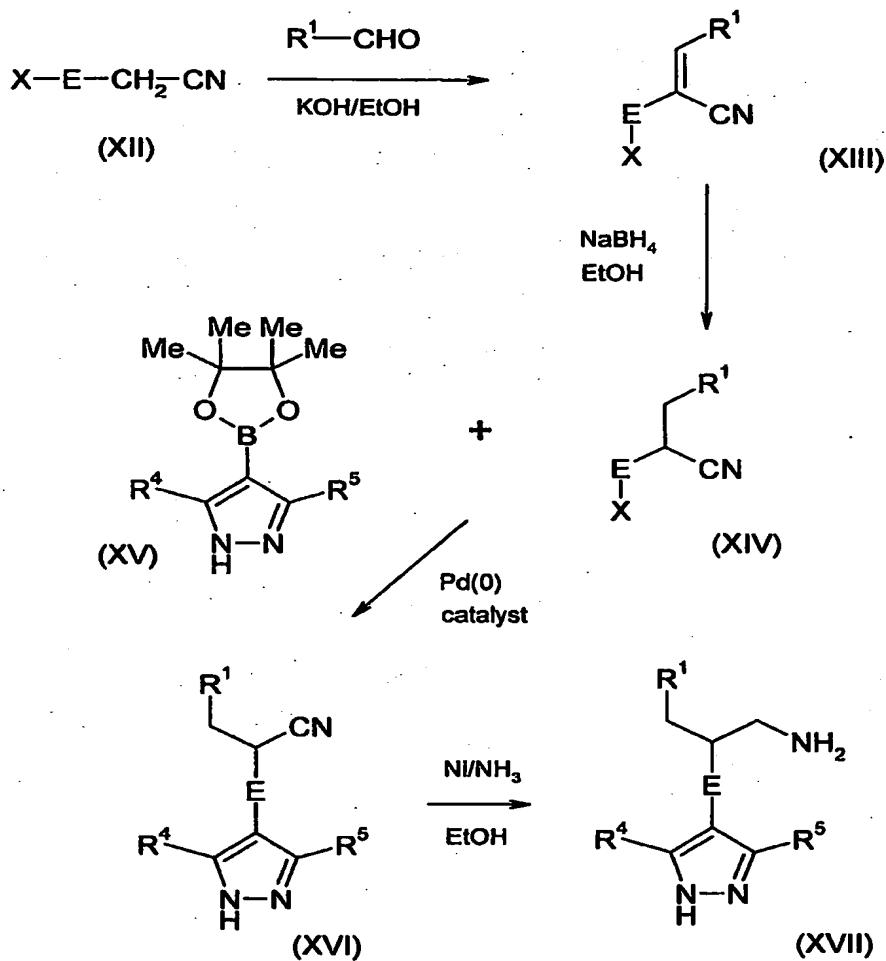
5 The reaction can be carried out under typical Suzuki Coupling conditions in the presence of a palladium catalyst such as bis(*tri-t-butylphosphine*)palladium and a base (e.g. a carbonate such as potassium carbonate). The reaction may be carried out in an aqueous solvent system, for example aqueous ethanol, and the reaction mixture is typically subjected to heating, for example to a temperature in excess of

10 100°C.

An illustrative synthetic route involving a Suzuki coupling step is shown in Scheme 1. The starting material for the synthetic route shown in scheme 1 is the halo-substituted aryl- or heteroaryl methyl nitrile (XII) in which X is a chlorine, bromine or iodine atom or a triflate group. The nitrile (XII) is condensed with the aldehyde 15 R<sup>1</sup>CHO in the presence of an alkali such as sodium or potassium hydroxide in an aqueous solvent system such as aqueous ethanol. The reaction can be carried out at room temperature.

The resulting substituted acrylonitrile derivative (XIII) is then treated with a reducing agent that will selectively reduce the alkene double bond without reducing 20 the nitrile group. A borohydride such as sodium borohydride may be used for this purpose to give the substituted acetonitrile derivative (XIV). The reduction reaction is typically carried out in a solvent such as ethanol and usually with heating, for example to a temperature up to about 65°C.

The reduced nitrile (XIV) is then coupled with the pyrazole boronate ester (XV) under the Suzuki coupling conditions described above to give a compound of the formula (I) in which A-NR<sup>2</sup>R<sup>3</sup> is a substituted acetonitrile group.



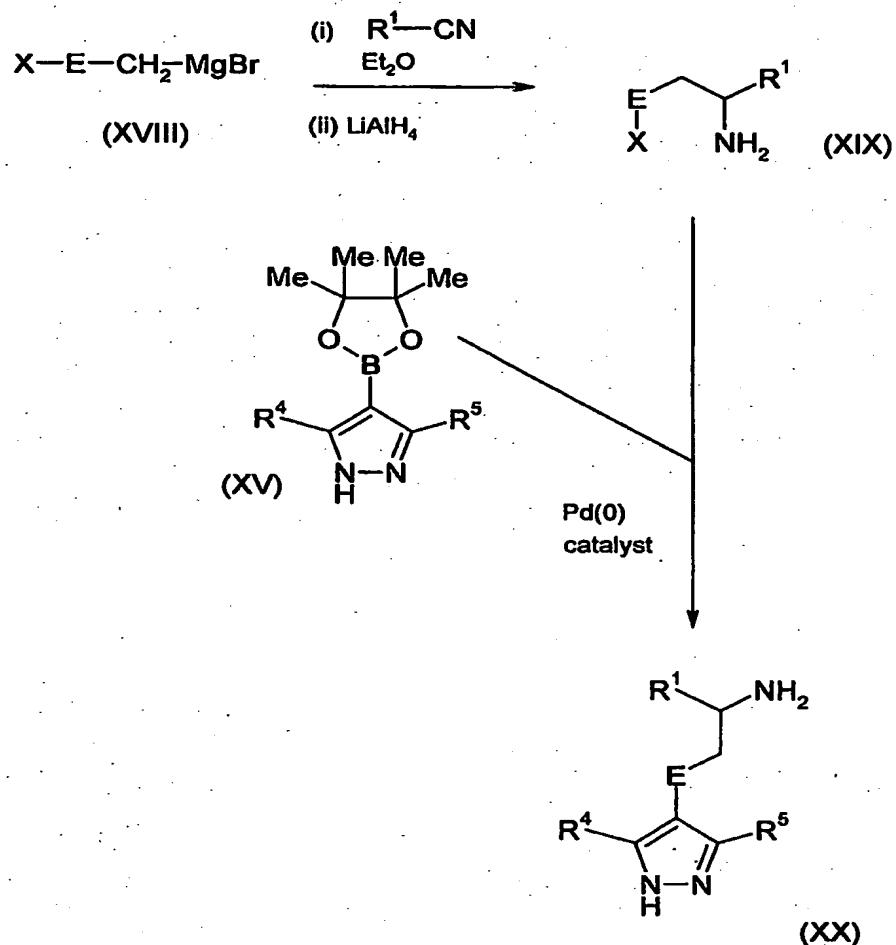
Scheme 1

5 The substituted acetonitrile compound (XVI) may then be reduced to the corresponding amine (XVII) by treatment with a suitable reducing agent such as Raney nickel and ammonia in ethanol.

The synthetic route shown in Scheme 1 gives rise to amino compounds of the formula (I) in which the aryl or heteroaryl group E is attached to the  $\beta$ -position of the group A relative to the amino group. In order to give amino compounds of the formula (I) in which  $R^1$  is attached to the  $\beta$ -position relative to the amino group, the 5 functional groups on the two starting materials in the condensation step can be reversed so that a compound of the formula  $X-E-CHO$  wherein X is bromine, chlorine, iodine or a triflate group is condensed with a compound of the formula  $R^1-CH_2-CN$  to give a substituted acrylonitrile derivative which is then reduced to the corresponding acetonitrile derivative before coupling with the pyrazole boronate 10 (XV) and reducing the cyano group to an amino group.

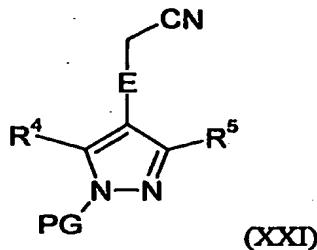
Compounds of the formula (I) in which  $R^1$  is attached to the  $\alpha$ -position relative to the amino group can be prepared by the sequence of reactions shown in Scheme 2.

In Scheme 2, the starting material is a halo-substituted aryl- or heteroarylmethyl 15 Grignard reagent (XVIII, X = bromine or chlorine) which is reacted with the nitrile  $R^1-CN$  in a dry ether such as diethyl ether to give an intermediate imine (not shown) which is reduced to give the amine (XIX) using a reducing agent such as lithium aluminium hydride. The amine (XIX) can be reacted with the boronate ester (XV) under the Suzuki coupling conditions described above to yield the amine (XX).



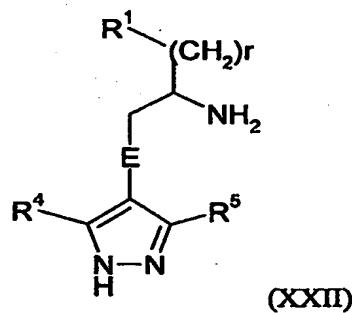
Scheme 2

Compounds of the formula (I) can also be prepared from the substituted nitrile compound (XXI):



wherein PG is a protecting group such as a tetrahydropyranyl group. The nitrile (XXI) can be condensed with an aldehyde of the formula  $R^1-(CH_2)_r-CHO$ , wherein r is 0 or 1, and the resulting substituted acrylonitrile subsequently reduced to the 5 corresponding substituted nitrile under conditions analogous to those set out in Scheme 1 above. The protecting group PG can then be removed by an appropriate method. The nitrile compound may subsequently be reduced to the corresponding amine by the use of a suitable reducing agent as described above.

The nitrile compound (XXI) may also be reacted with a Grignard reagent of the 10 formula  $R^1-(CH_2)_r-MgBr$  under standard Grignard reaction conditions followed by deprotection to give an amino compound of the invention which has the structure shown in formula (XXII).



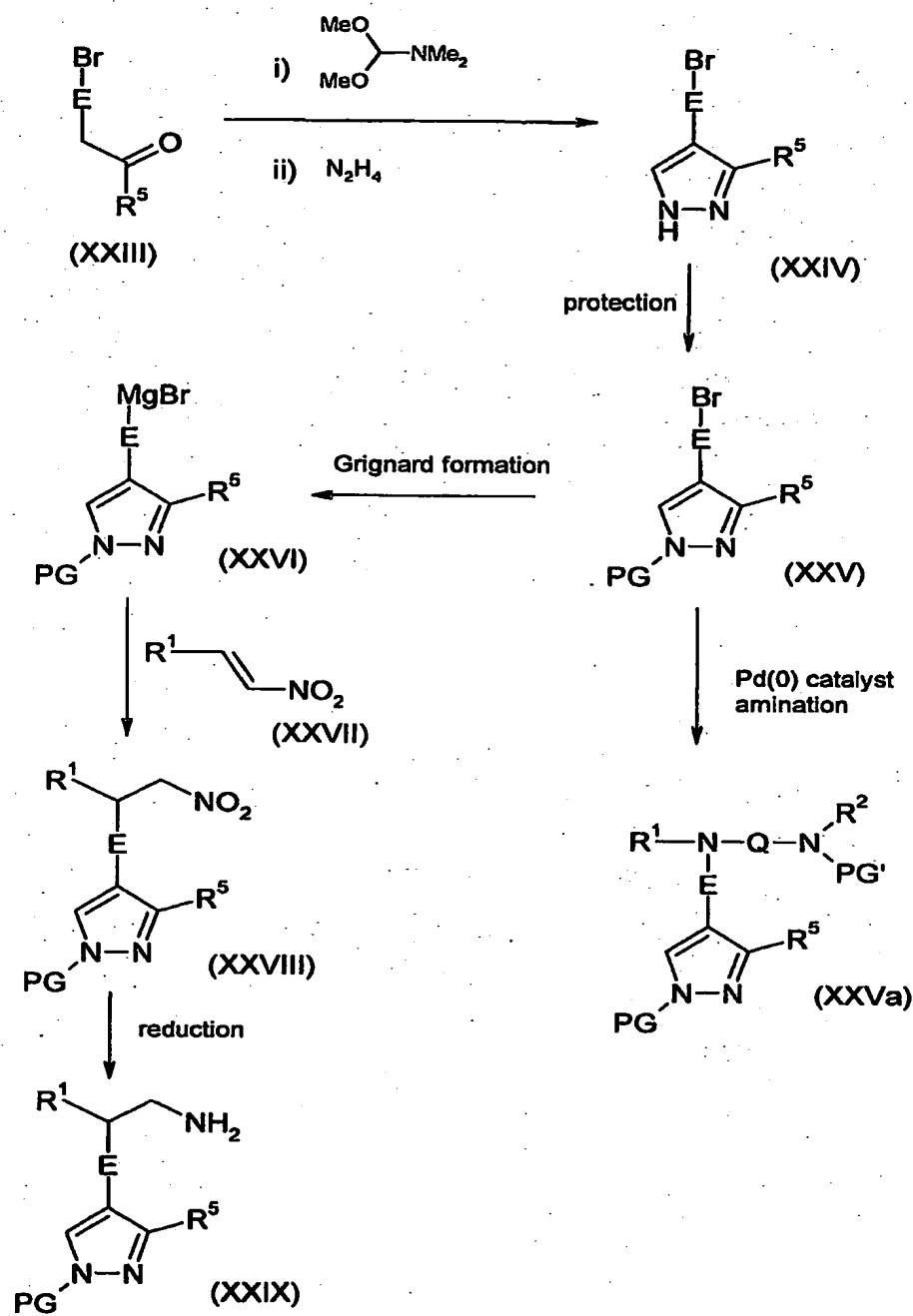
In the preparative procedures outlined above, the coupling of the aryl or heteroaryl 15 group E to the pyrazole is accomplished by reacting a halo-pyrazole or halo-aryl or heteroaryl compound with a boronate ester or boronic acid in the presence of a palladium catalyst and base. Many boronates suitable for use in preparing compounds of the invention are commercially available, for example from Boron

Molecular Limited of Noble Park, Australia, or from Combi-Blocks Inc, of San Diego, USA. Where the boronates are not commercially available, they can be prepared by methods known in the art, for example as described in the review article by N. Miyaura and A. Suzuki, *Chem. Rev.* 1995, 95, 2457. Thus, boronates 5 can be prepared by reacting the corresponding bromo-compound with an alkyl lithium such as butyl lithium and then reacting with a borate ester. The resulting boronate ester derivative can, if desired, be hydrolysed to give the corresponding boronic acid.

Another synthetic route to compounds of the formula (I) is shown in Scheme 3.

10 In Scheme 3, a pyrazole ring system is constructed by reacting a halogenated arylmethyl ketone (XXIII) with dimethylformamide acetal followed by hydrazine. The reaction typically proceeds in two steps. The reaction between the acetal and the ketone is brought about by heating to an elevated temperature (e.g. 80-100°C) for a period of several hours to give an intermediate (not shown) which is then 15 treated with hydrazine to give the bromo-aryl-pyrazole (XXIV). The reaction with hydrazine is typically carried out in a polar solvent such as ethanol at an elevated temperature, for example under conditions of reflux.

P033 US



Scheme 3

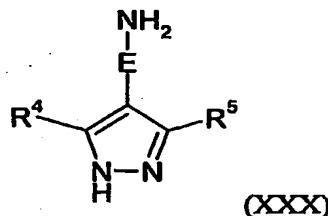
P033 US

The presence of the bromine atom in the bromo-aryl-pyrazole (XXIV) enables it to be used in a number of different synthetic procedures leading to compounds of the formula (I). In one procedure, the pyrazole 1-(H) position may first be protected, for example by the introduction of a protecting group (PG) such as a tetrahydro-5 pyranyl group or a 4-methoxybenzyl group, to give a protected derivative (XXV). The protected bromo-aryl-pyrazole can then be converted into an organometallic reagent such as a Grignard reagent (XXVI) by methods (e.g. by reaction with magnesium in an ether solvent) that are well known to those skilled in the art of organic chemistry. The Grignard reagent (XXVI) can be reacted with a variety of 10 substrates to give compounds of the formula (I) or precursors thereto. For example, reaction of the Grignard reagent (XXVI) with an aryl nitroalkene (XXVII) such as nitrostyrene gives rise to the substituted nitroethane derivative (XXVIII) which can be reduced (for example by hydrogenation over a Pd/C or Raney nickel catalyst, or by reaction with LiAlH<sub>4</sub> or sodium bis-(2-methoxyethoxy) aluminium dihydride) to 15 give an amine (XXIX) of the invention. The aryl-nitroalkene (XXVII) can be prepared by the well known Knoevenagel condensation of an aryl aldehyde with nitromethane.

Compounds of the formula (I) in which the group A contains a nitrogen atom attached to the group E can be prepared from the protected bromo-aryl-pyrazole 20 (XXV) by palladium catalysed amination using an amine of the formula R<sup>1</sup>-NH-Q-NR<sup>2</sup>-PG', where Q is an alkylene residue of the group A, and PG' is a protecting group such as a butyloxycarbonyl group, under palladium catalysed amination conditions of the type described in *Organic Letters*, 2002, vol. 4, No. 17, pp2885-2888. The resulting compound of the formula (XXVa) can be deprotected by 25 removal of the protecting groups using standard methods to give the compound of the formula (I).

Compounds of the formula (I) in which the group A contains a nitrogen atom attached to the group E can also be prepared by well known synthetic procedures from compounds of the formula (XXX) or a protected form thereof. Compounds of 30 the formula (XXX) can be obtained by a Suzuki coupling reaction of a compound

of the formula (XV) (see Scheme 1) with a compound of the formula Br-E-NH<sub>2</sub> such as 4-bromoaniline.



Once formed, many compounds of the formula (I) can be converted into other 5 compounds of the formula (I) using standard functional group interconversions. For example, compounds of the formula (I) in which the NR<sup>2</sup>R<sup>3</sup> forms part of a nitrile group can be reduced to the corresponding amine. Compounds in which NR<sup>2</sup>R<sup>3</sup> is an NH<sub>2</sub> group can be converted to the corresponding alkylamine by reductive alkylation, or to a cyclic group. Examples of functional group 10 interconversions and reagents and conditions for carrying out such conversions can be found in, for example, *Advanced Organic Chemistry*, by Jerry March, 4<sup>th</sup> edition, 119, Wiley Interscience, New York, *Fiesers' Reagents for Organic Synthesis*, Volumes 1-17, John Wiley, edited by Mary Fieser (ISBN: 0-471-58283-2), and *Organic Syntheses*, Volumes 1-8, John Wiley, edited by Jeremiah P. Freeman 15 (ISBN: 0-471-31192-8).

In many of the reactions described above, it may be necessary to protect one or more groups to prevent reaction from taking place at an undesirable location on the molecule. Examples of protecting groups, and methods of protecting and deprotecting functional groups, can be found in *Protective Groups in Organic 20 Synthesis* (T. Green and P. Wuts; 3rd Edition; John Wiley and Sons, 1999).

A hydroxy group may be protected, for example, as an ether (-OR) or an ester (-OC(=O)R), for example, as: a t-butyl ether; a benzyl, benzhydryl (diphenylmethyl), or trityl (triphenylmethyl) ether; a trimethylsilyl or t-butyldimethylsilyl ether; or an acetyl ester (-OC(=O)CH<sub>3</sub>, -OAc). An aldehyde or ketone group may be protected, 25 for example, as an acetal (R-CH(OR)<sub>2</sub>) or ketal (R<sub>2</sub>C(OR)<sub>2</sub>), respectively, in which

the carbonyl group (>C=O) is converted to a diether (>C(OR)<sub>2</sub>), by reaction with, for example, a primary alcohol. The aldehyde or ketone group is readily regenerated by hydrolysis using a large excess of water in the presence of acid. An amine group may be protected, for example, as an amide (-NRCO-R) or a urethane

5 (-NRCO-OR), for example, as: a methyl amide (-NHCO-CH<sub>3</sub>); a benzyloxy amide (-NHCO-OCH<sub>2</sub>C<sub>6</sub>H<sub>5</sub>, -NH-Cbz); as a t-butoxy amide (-NHCO-OC(CH<sub>3</sub>)<sub>3</sub>, -NH-Boc); a 2-biphenyl-2-propoxy amide (-NHCO-OC(CH<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>4</sub>C<sub>6</sub>H<sub>5</sub>, -NH-Bpoc), as a 9-fluorenylmethoxy amide (-NH-Fmoc), as a 6-nitroveratryloxy amide (-NH-Nvoc), as a 2-trimethylsilylethoxy amide (-NH-Teoc), as a 2,2,2-

10 trichloroethoxy amide (-NH-Troc), as an allyloxy amide (-NH-Alloc), or as a 2-(phenylsulphonyl)ethoxy amide (-NH-Psec). Other protecting groups for amines, such as cyclic amines and heterocyclic N-H groups, include toluenesulphonyl (tosyl) and methanesulphonyl (mesyl) groups and benzyl groups such as a *para*-methoxybenzyl (PMB) group. A carboxylic acid group may be

15 protected as an ester for example, as: an C<sub>1-7</sub> alkyl ester (e.g., a methyl ester; a t-butyl ester); a C<sub>1-7</sub> haloalkyl ester (e.g., a C<sub>1-7</sub> trihaloalkyl ester); a triC<sub>1-7</sub> alkylsilyl-C<sub>1-7</sub>alkyl ester; or a C<sub>5-20</sub> aryl-C<sub>1-7</sub> alkyl ester (e.g., a benzyl ester; a nitrobenzyl ester); or as an amide, for example, as a methyl amide. A thiol group may be protected, for example, as a thioether (-SR), for example, as: a benzyl thioether; an

20 acetamidomethyl ether (-S-CH<sub>2</sub>NHC(=O)CH<sub>3</sub>).

The 1(H) position of the pyrazole group in the compounds of the formula (I) or its precursors can be protected by a variety of groups, the protecting group being selected according to the nature of the reaction conditions to which the group is exposed. Examples of protecting groups for the pyrazole N-H include

25 tetrahydropyranyl, benzyl and 4-methoxybenzyl groups.

Many of the chemical intermediates described above are novel and such novel intermediates form a further aspect of the invention.

#### Pharmaceutical Formulations

The invention also provides compounds of the formula (I) as hereinbefore defined in the form of pharmaceutical compositions.

The pharmaceutical compositions can be in any form suitable for oral, parenteral, topical, intranasal, ophthalmic, otic, rectal, intra-vaginal, or transdermal

5 administration. Where the compositions are intended for parenteral administration, they can be formulated for intravenous, intramuscular, intraperitoneal, or subcutaneous administration or for direct delivery into a target organ or tissue by injection, infusion or other means of delivery.

10 Pharmaceutical dosage forms suitable for oral administration include tablets, capsules, caplets, pills, lozenges, syrups, solutions, powders, granules, elixirs and suspensions, sublingual tablets, wafers or patches and buccal patches.

15 Pharmaceutical compositions containing compounds of the formula (I) can be formulated in accordance with known techniques, see for example, Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, PA, USA.

20 Thus, tablet compositions can contain a unit dosage of active compound together with an inert diluent or carrier such as a sugar or sugar alcohol, eg; lactose, sucrose, sorbitol or mannitol; and/or a non-sugar derived diluent such as sodium carbonate, calcium phosphate, calcium carbonate, or a cellulose or derivative thereof such as methyl cellulose, ethyl cellulose, hydroxypropyl methyl cellulose, and starches such as corn starch. Tablets may also contain such standard ingredients as binding and granulating agents such as polyvinylpyrrolidone, disintegrants (e.g. swellable crosslinked polymers such as crosslinked carboxymethylcellulose), lubricating agents (e.g. stearates), preservatives (e.g. parabens), antioxidants (e.g. BHT), buffering agents (for example phosphate or citrate buffers), and effervescent agents such as citrate/bicarbonate mixtures. Such excipients are well known and do not need to be discussed in detail here.

Capsule formulations may be of the hard gelatin or soft gelatin variety and can contain the active component in solid, semi-solid, or liquid form. Gelatin capsules can be formed from animal gelatin or synthetic or plant derived equivalents thereof.

The solid dosage forms (e.g. tablets, capsules etc.) can be coated or un-coated, but

5 typically have a coating, for example a protective film coating (e.g. a wax or varnish) or a release controlling coating. The coating (e.g. a Eudragit <sup>TM</sup> type polymer) can be designed to release the active component at a desired location within the gastro-intestinal tract. Thus, the coating can be selected so as to degrade under certain pH conditions within the gastrointestinal tract, thereby selectively

10 release the compound in the stomach or in the ileum or duodenum.

Instead of, or in addition to, a coating, the drug can be presented in a solid matrix comprising a release controlling agent, for example a release delaying agent which may be adapted to selectively release the compound under conditions of varying acidity or alkalinity in the gastrointestinal tract. Alternatively, the matrix material

15 or release retarding coating can take the form of an erodible polymer (e.g. a maleic anhydride polymer) which is substantially continuously eroded as the dosage form passes through the gastrointestinal tract.

Compositions for topical use include ointments, creams, sprays, patches, gels, liquid drops and inserts (for example intraocular inserts). Such compositions can be

20 formulated in accordance with known methods.

Compositions for parenteral administration are typically presented as sterile aqueous or oily solutions or fine suspensions, or may be provided in finely divided sterile powder form for making up extemporaneously with sterile water for injection.

25 Examples of formulations for rectal or intra-vaginal administration include pessaries and suppositories which may be, for example, formed from a shaped mouldable or waxy material containing the active compound.

Compositions for administration by inhalation may take the form of inhalable powder compositions or liquid or powder sprays, and can be administrated in standard form using powder inhaler devices or aerosol dispensing devices. Such devices are well known. For administration by inhalation, the powdered

5 formulations typically comprise the active compound together with an inert solid powdered diluent such as lactose.

The compounds of the inventions will generally be presented in unit dosage form and, as such, will typically contain sufficient compound to provide a desired level of biological activity. For example, a formulation intended for oral administration

10 may contain from 0.1 milligrams to 2 grams of active ingredient, more usually from 10 milligrams to 1 gram, for example, 50 milligrams to 500 milligrams.

The active compound will be administered to a patient in need thereof (for example a human or animal patient) in an amount sufficient to achieve the desired therapeutic effect.

15 **Protein Kinase Inhibitory Activity**

The activity of the compounds of the invention as inhibitors of protein kinases A and B can be measured using the assays set forth in the examples below and the level of activity exhibited by a given compound can be defined in terms of the IC<sub>50</sub> value. Preferred compounds of the present invention are compounds having an IC<sub>50</sub>

20 value of less than 1 micromole, more preferably less than 0.1 micromole.

**Therapeutic Uses**

**Prevention or Treatment of Proliferative Disorders**

The compounds of the formula (I) are inhibitors of protein kinase A and protein kinase B. As such, they are expected to be useful in providing a means of

25 preventing the growth or inducing apoptosis of neoplasias. It is therefore anticipated that the compounds will prove useful in treating or preventing proliferative disorders such as cancers. In particular tumours with deletions or

inactivating mutations in PTEN may be particularly sensitive to PKB inhibitors. Tumours which have other abnormalities leading to an upregulated PKB pathway signal may also be particularly sensitive to inhibitors of PKB. Examples of such abnormalities include but are not limited to overexpression of one or more PI3K

5 subunits, over-expression of one or more PKB isoforms, or mutations in PI3K, PDK1, or PKB which lead to an increase in the basal activity of the enzyme in question.

It is also envisaged that the compounds of the invention will be useful in treating other conditions which result from disorders in proliferation or survival such as

10 viral infections, and neurodegenerative diseases for example. PKB plays an important role in maintaining the survival of immune cells during an immune response and therefore PKB inhibitors could be particularly beneficial in immune disorders including autoimmune conditions.

Therefore, PKB inhibitors could be useful in the treatment of diseases in which 15 there is a disorder of proliferation, apoptosis or differentiation.

Examples of cancers which may be inhibited include, but are not limited to, a carcinoma, for example a carcinoma of the bladder, breast, colon (e.g. colorectal carcinomas such as colon adenocarcinoma and colon adenoma), kidney, epidermal, liver, lung, for example adenocarcinoma, small cell lung cancer and non-small cell 20 lung carcinomas, oesophagus, gall bladder, ovary, pancreas e.g. exocrine pancreatic carcinoma, stomach, cervix, thyroid, prostate, or skin, for example squamous cell carcinoma; a hematopoietic tumour of lymphoid lineage, for example leukaemia, acute lymphocytic leukaemia, B-cell lymphoma, T-cell lymphoma, Hodgkin's lymphoma, non-Hodgkin's lymphoma, hairy cell lymphoma, or Burkett's 25 lymphoma; a hematopoietic tumour of myeloid lineage, for example acute and chronic myelogenous leukaemias, myelodysplastic syndrome, or promyelocytic leukaemia; thyroid follicular cancer; a tumour of mesenchymal origin, for example fibrosarcoma or habdomyosarcoma; a tumour of the central or peripheral nervous system, for example astrocytoma, neuroblastoma, glioma or schwannoma;

melanoma; seminoma; teratocarcinoma; osteosarcoma; xenoderoma pigmentosum; keratoctanthoma; thyroid follicular cancer; or Kaposi's sarcoma.

Thus, in the pharmaceutical compositions, uses or methods of this invention for treating a disease or condition comprising abnormal cell growth, the disease or 5 condition comprising abnormal cell growth in one embodiment is a cancer.

Particular subsets of cancers include breast cancer, ovarian cancer, colon cancer, prostate cancer, oesophageal cancer, squamous cancer and non-small cell lung carcinomas.

It is also possible that some protein kinase B inhibitors can be used in combination 10 with other anticancer agents. For example, it may be beneficial to combine of an inhibitor that induces apoptosis with another agent which acts via a different mechanism to regulate cell growth thus treating two of the characteristic features of cancer development. Examples of such combinations are set out below.

#### Immune Disorders

15 Immune disorders for which PKA and PKB inhibitors may be beneficial include but are not limited to autoimmune conditions and chronic inflammatory diseases, for example systemic lupus erythematosus, autoimmune mediated glomerulonephritis, rheumatoid arthritis, psoriasis, inflammatory bowel disease, and autoimmune diabetes mellitus, Eczema hypersensitivity reactions, asthma, COPD, rhinitis, and 20 upper respiratory tract disease.

#### Other Therapeutic Uses

25 PKB plays a role in apoptosis, proliferation, differentiation and therefore PKB inhibitors could also be useful in the treatment of the following diseases other than cancer and those associated with immune dysfunction; viral infections, for example herpes virus, pox virus, Epstein-Barr virus, Sindbis virus, adenovirus, HIV, HPV, HCV and HCMV; prevention of AIDS development in HIV-infected individuals; cardiovascular diseases for example cardiac hypertrophy, restenosis,

atherosclerosis; neurodegenerative disorders, for example Alzheimer's disease, AIDS-related dementia, Parkinson's disease, amyotrophic lateral sclerosis, retinitis pigmentosa, spinal muscular atrophy and cerebellar degeneration; glomerulonephritis; myelodysplastic syndromes, ischemic injury associated 5 myocardial infarctions, stroke and reperfusion injury, degenerative diseases of the musculoskeletal system, for example, osteoporosis and arthritis, aspirin-sensitive rhinosinusitis, cystic fibrosis, multiple sclerosis, kidney diseases.

#### Methods of Treatment

It is envisaged that the compounds of the formula (I) will be useful in the prophylaxis 10 or treatment of a range of disease states or conditions mediated by protein kinase A and/or protein kinase B. Examples of such disease states and conditions are set out above.

Compounds of the formula (I) are generally administered to a subject in need of such administration, for example a human or animal patient, preferably a human. 15 The compounds will typically be administered in amounts that are therapeutically or prophylactically useful and which generally are non-toxic. However, in certain situations (for example in the case of life threatening diseases), the benefits of administering a compound of the formula (I) may outweigh the disadvantages of any toxic effects or side effects, in which case it may be considered desirable to 20 administer compounds in amounts that are associated with a degree of toxicity.

The compounds may be administered over a prolonged term to maintain beneficial therapeutic effects or may be administered for a short period only. Alternatively they may be administered in a pulsatile manner.

A typical daily dose of the compound can be in the range from 100 picograms to 25 100 milligrams per kilogram of body weight, more typically 10 nanograms to 10 milligrams per kilogram of bodyweight although higher or lower doses may be administered where required. Ultimately, the quantity of compound administered

will be commensurate with the nature of the disease or physiological condition being treated and will be at the discretion of the physician.

The compounds of the formula (I) can be administered as the sole therapeutic agent or they can be administered in combination therapy with one or more other

5 compounds for treatment of a particular disease state, for example a neoplastic disease such as a cancer as hereinbefore defined. Examples of other therapeutic agents that may be administered together (whether concurrently or at different time intervals) with the compounds of the formula (I) include but are not limited to topoisomerase inhibitors, alkylating agents, antimetabolites, DNA binders and

10 microtubule inhibitors, such as cisplatin, cyclophosphamide, doxorubicin, irinotecan, fludarabine, 5FU, taxanes, mitomycin C or radiotherapy. For the case of protein kinase B inhibitors combined with other therapies the two or more treatments may be given in individually varying dose schedules and via different routes.

15 **EXPERIMENTAL**

The invention will now be illustrated, but not limited, by reference to the specific embodiments described in the following procedures and examples.

The starting materials for each of the procedures described below are commercially available unless otherwise specified.

20 In the examples, the compounds prepared were characterised by liquid chromatography and mass spectroscopy using the system and operating conditions set out below. Where chlorine is present, the mass quoted for the compound is for  $^{35}\text{Cl}$ . The two systems were equipped with identical chromatography columns and were set up to run under the same operating conditions. The operating conditions

25 used are also described below.

**Platform System**

HPLC System: Waters 2795

Mass Spec Detector: Micromass Platform LC

P033 US

PDA Detector: Waters 2996 PDA

Acidic Analytical conditions:

Eluent A: H<sub>2</sub>O (0.1% Formic Acid)

Eluent B: CH<sub>3</sub>CN (0.1% Formic Acid)

5 Gradient: 5-95% eluent B over 3.5 minutes

Flow: 1.5 ml/min

Column: Phenomenex Synergi 4μ Max-RP 80A, 50x4.6mm

Basic Analytical conditions:

Eluent A: H<sub>2</sub>O (10mM NH<sub>4</sub>HCO<sub>3</sub> buffer adjusted to pH=9.5 with NH<sub>4</sub>OH)

10 Eluent B: CH<sub>3</sub>CN

Gradient: 05-95% eluent B over 3.5 minutes

Flow: 1.5 ml/min

Column: Waters Xterra MS C<sub>18</sub> 5μm 4.6x50mm

Polar Analytical conditions:

15 Eluent A: H<sub>2</sub>O (0.1% Formic Acid)

Eluent B: CH<sub>3</sub>CN (0.1% Formic Acid)

Gradient: 00-50% eluent B over 3 minutes

Flow: 1.5 ml/min

Column: Phenomenex Synergi 4μ Hydro 80A, 50x4.6mm

20 MS conditions:

Capillary voltage: 3.5 kV

Cone voltage: 30 V

Source Temperature: 120 °C

Scan Range: 165-700 amu

25 Ionisation Mode: Electrospray Negative, Positive or Positive & Negative

FractionLynx System

System: Waters FractionLynx (dual analytical/prep)

P033 US

HPLC Pump: Waters 2525  
Injector-Autosampler: Waters 2767  
Mass Spec Detector: Waters-Micromass ZQ  
PDA Detector: Waters 2996 PDA

5 Acidic Analytical conditions:

Eluent A: H<sub>2</sub>O (0.1% Formic Acid)  
Eluent B: CH<sub>3</sub>CN (0.1% Formic Acid)  
Gradient: 5-95% eluent B over 5 minutes  
Flow: 2.0 ml/min  
10 Column: Phenomenex Synergi 4μ Max-RP 80A, 50x4.6mm

Polar Analytical conditions:

Eluent A: H<sub>2</sub>O (0.1% Formic Acid)  
Eluent B: CH<sub>3</sub>CN (0.1% Formic Acid)  
Gradient: 00-50% eluent B over 5 minutes  
15 Flow: 2.0 ml/min  
Column: Phenomenex Synergi 4μ Max-RP 80A, 50x4.6mm

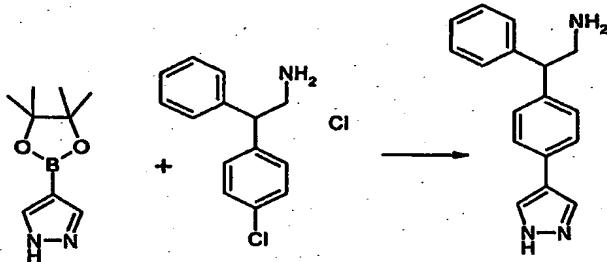
MS conditions:

Capillary voltage: 3.5 kV  
Cone voltage: 25 V  
20 Source Temperature: 120 °C  
Scan Range: 125-800 amu  
Ionisation Mode: ElectroSpray Positive or ElectroSpray Positive & Negative

In the examples below, the following key is used to identify the LCMS conditions used:

25 PS-A Platform System – acidic analytical conditions  
PS-B Platform System – basic analytical conditions  
PS-P Platform System – polar analytical conditions  
FL-A FractionLynx System – acidic analytical conditions  
FL-P FractionLynx System – polar analytical conditions

P033 US

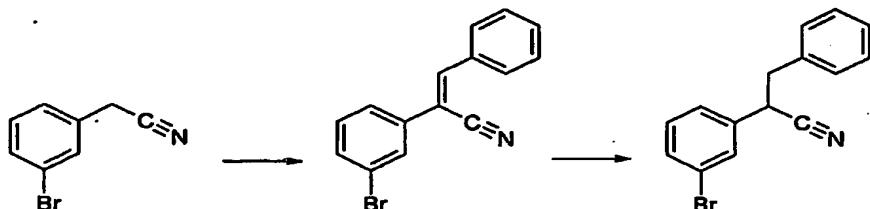
EXAMPLE 12-Phenyl-2-[4-(1H-pyrazol-4-yl)-phenyl]-ethylamine

5 To a suspension of 2-(4-chlorophenyl)-2-phenylethylamine hydrochloride (134 mg, 0.5 mmol, 1.0 equiv.) (Array PPA-Q02-1) in toluene (0.8 ml) was added bis(tri-*t*-butylphosphine)palladium (0) (3 mg, 1 mol%) (Strem) and the mixture was purged with nitrogen. A suspension of 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1H-pyrazole (107 mg, 0.55 mmol, 1.1 equiv.) (Aldrich 52,505-7) in ethanol (0.8 ml)

10 was added followed by potassium carbonate (415 mg, 3.0 mmol, 6 equiv.) in water (2.5 ml). The mixture was purged with nitrogen and sealed. The reaction mixture was heated in a CEM Explorer™ microwave to 135 °C for 15 minutes using 50 watts power. The solvents were removed and the residue was partitioned between ethyl acetate and 2N NaOH. The aqueous layer was extracted with ethyl acetate and the combined organic layers were washed with brine, dried ( $\text{MgSO}_4$ ) and concentrated under reduced pressure. The crude reaction mixture was purified by column chromatography ( $\text{SiO}_2$ ), eluting with a mixture of dichloromethane (90ml): methanol (18ml): acetic acid (3ml):  $\text{H}_2\text{O}$  (2ml) to afford the title compound 14 mg (9%); LCMS (PS-A)  $R_t$  1.79 min;  $m/z$   $[\text{M}+\text{H}]^+$  264.

15

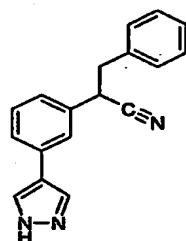
EXAMPLE 23-Phenyl-2-[3-(1H-pyrazol-4-yl)-phenyl]-propionitrile2A. 2-(3-Bromo-phenyl)-3-phenyl-propionitrile



A solution of 40% KOH (2.83 g in 5.0 ml of H<sub>2</sub>O) in ethanol (13 ml) was added to a solution of benzaldehyde (2.85 ml, 28.05 mmol) and 3-bromophenylacetonitrile (5 g, 25.50 mmol) in ethanol (9 ml). The reaction mixture was then stirred at room 5 temperature for 2 hours and the precipitate was collected by suction filtration and washed with cold ethanol (6.68 g, 92 %). The crude product (3.45g, 12.14 mmol) was then dissolved in ethanol (35 ml) and heated to 65 °C. Sodium borohydride (459 mg, 12.14 mmol) was added in portions and the reaction mixture was maintained at this temperature for a further 2 hours. Upon cooling, water (10 ml) 10 was added and the solvent was removed under reduced pressure. The residue was partitioned between water (100 ml) and ethyl acetate (100 ml). The organic layer was separated, dried (MgSO<sub>4</sub>), filtered and concentrated to afford the desired product (1.80 g, 52 %), which was used without purification.

**2B. 3-Phenyl-2-[3-(1H-pyrazol-4-yl)-phenyl]-propionitrile**

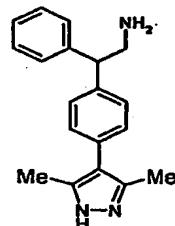
15



2-(3-Bromo-phenyl)-3-phenyl-propionitrile was reacted with 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1H-pyrazole following the procedure set out in Example 1 to give the title compound. (LC/MS: (PS-A) R<sub>t</sub> 2.98 [M+H]<sup>+</sup> 274).

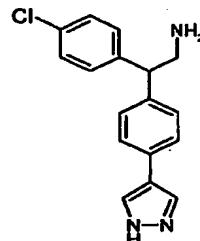
20 **EXAMPLE 3**

P033 US

2-[4-(3,5-Dimethyl-1H-pyrazol-4-yl)-phenyl]-2-phenyl-ethylamine

Following the procedure of Example 1 but using 3,5-dimethyl-4-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-1H-pyrazole (Boron Molecular D03-BM152)

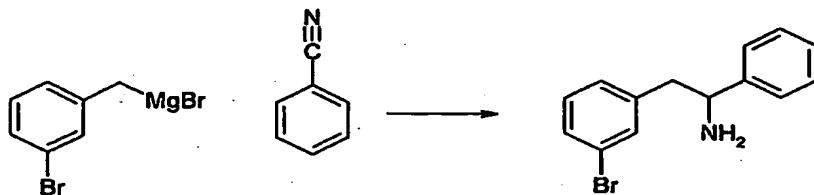
5 instead of 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1H-pyrazole gave the title compound. (LC/MS: (PS-A) R<sub>t</sub> 1.79 [M+H]<sup>+</sup> 292.

EXAMPLE 42-(4-Chloro-phenyl)-2-[4-(1H-pyrazol-4-yl)-phenyl]-ethylamine

10 Following the procedure of Example 1 but using 2,2-bis-(4-chlorophenyl)-ethylamine in place of 2-(4-chlorophenyl)-2-phenylethylamine hydrochloride\* gave the title compound. (LC/MS: (PS-A) R<sub>t</sub> 1.99 [M+H]<sup>+</sup> 298).

\*This starting material can be made by the method described in *J. Amer. Chem. Soc.*, 1983, 105, 3183-3188.

EXAMPLE 52-[3-(3,5-Dimethyl-1H-pyrazol-4-yl)-phenyl]-1-phenyl-ethylamine5A. 2-(3-Bromo-phenyl)-1-phenyl-ethylamine

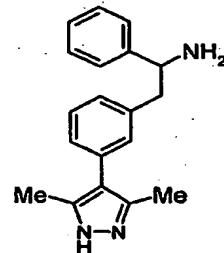


Benzonitrile (500 mg, 4.849 mmol) was added dropwise to a solution of 3-bromobenzylmagnesium bromide (0.275 M solution in diethyl ether, 21.1 ml, 5.818 mmol) under an atmosphere of nitrogen at room temperature. The reaction mixture

5 was then heated to reflux for a period of 2 hours then allowed to cool. Lithium aluminium hydride (1.0 M in THF, 4.85 ml, 4.849 mmol) was then added cautiously and the reaction mixture was allowed to heat at reflux for a further 16 hours. Upon cooling, the reaction was quenched by cautious and dropwise addition of water (5 ml) and then partitioned between water (20 ml) and ethyl acetate (100 ml). The organic layer was separated, dried ( $\text{MgSO}_4$ ), filtered and concentrated.

10 Purification by ion exchange chromatography afforded the desired compound (420 mg, 31 %).

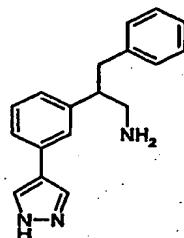
**5B. 2-[3-(3,5-Dimethyl-1H-pyrazol-4-yl)-phenyl]-1-phenyl-ethylamine**



15 The product of 5B was reacted with 3,5-dimethyl-4-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-1H-pyrazole following the procedure set out in Example 1 to give the title compound. (LC/MS: (PS-B)  $R_t$  2.54  $[\text{M}+\text{H}]^+$  292).

**EXAMPLE 6**

**3-Phenyl-2-[3-(1H-pyrazol-4-yl)-phenyl]-propylamine**

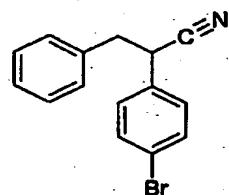


To a solution of the product of Example 2 (70 mg, 0.256 mmol, 1.0 equiv) in ethanol (25 ml) was added concentrated ammonia (0.5 ml) and Raney Nickel (approximately 0.5 ml of the water suspension) and the reaction mixture was subjected to a hydrogen atmosphere for 17 hours. The mixture was filtered through Celite® and the mother liquor was concentrated under reduced pressure to give the title compound which was purified by preparative liquid chromatography. (LC/MS: (PS-A)  $R_t$  1.89  $[M+H]^+$  278.

#### EXAMPLE 7

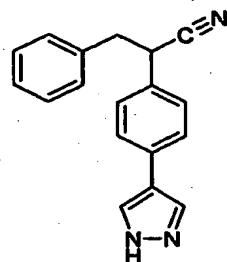
**10 3-Phenyl-2-[4-(1H-pyrazol-4-yl)-phenyl]-propylamine**

**7A. 2-(4-Bromo-phenyl)-3-phenyl-propionitrile**



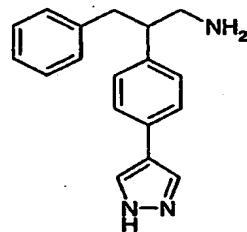
Following the procedure described in Example 2A but substituting 4-bromophenylacetonitrile for 3-bromophenylacetonitrile gave the title compound 15 was obtained which was used in the next step without further purification.

**7B. 3-Phenyl-2-[4-(1H-pyrazol-4-yl)-phenyl]-propionitrile**



By following the procedure described in Example 1 but substituting 2-(4-Bromo-phenyl)-3-phenyl-propionitrile for 2-(4-chlorophenyl)-2-phenylethylamine, the title compound was obtained.

5 7C. 3-Phenyl-2-[4-(1H-pyrazol-4-yl)-phenyl]-propylamine

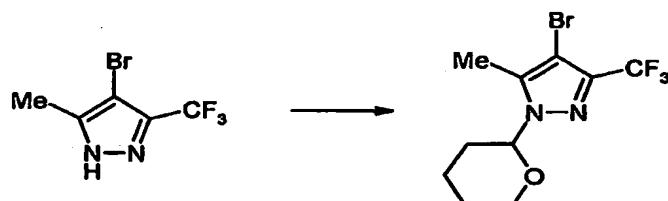


The nitrile product of Example 7B was reduced using the conditions described in Example 6 to give the title compound. (LC/MS: (PS-B)  $R_f$  3.03  $[M+H]^+$  278.

EXAMPLE 8

10 [4-(5-Methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-acetonitrile

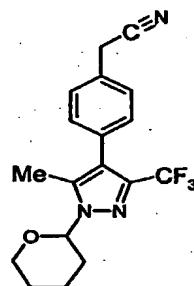
8A. 4-Bromo-5-methyl-1-(tetrahydro-pyran-2-yl)-3-trifluoromethyl-1H-pyrazole



15 To a solution of 4-bromo-5-methyl-3-trifluoromethyl-1H-pyrazole (1.4 g, 6.2 mmol, 1.0 equiv) in chloroform (31 ml) was added p-toluene sulphonic acid monohydrate (118 mg, 0.62 mmol, 0.1 equiv). The solution was cooled to 0 °C and

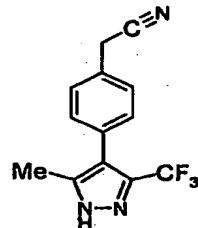
3,4-dihydro-2H-pyran (0.85 ml, 9.3 mmol, 1.5 equiv) was added drop-wise over 5 minutes. The mixture was allowed to warm to room temperature for 1 hour and the solvents were removed under reduced pressure. The crude mixture was purified by column chromatography (SiO<sub>2</sub>), eluting with 0→25% EtOAc-petrol over a linear gradient to afford the title compound 1.4 g (59%), LCMS (PS-A) R<sub>t</sub> 3.72 min; *m/z* [M+H]<sup>+</sup> 314.

**8B. {4-[5-Methyl-1-(tetrahydro-pyran-2-yl)-3-trifluoromethyl-1H-pyrazol-4-yl]-phenyl}-acetonitrile**



10 The product of Example 8A, 4-bromo-5-methyl-1-(tetrahydro-pyran-2-yl)-3-trifluoromethyl-1H-pyrazole, was reacted with 4-(cyanomethylphenyl)boronic acid (Combi-Blocks, San Diego, USA Cat. No. 2444-001), under the conditions described in Example 1, to give the title compound.

**8C. [4-(5-Methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-acetonitrile**



15 To {4-[5-Methyl-1-(tetrahydro-pyran-2-yl)-3-trifluoromethyl-1H-pyrazol-4-yl]-phenyl}-acetonitrile (Example 8B) (35 mg, 0.1 mmol, 1.0 equiv) in ethyl acetate (1 ml) was added HCl in ethyl acetate (1 ml) and the mixture was stirred for 1 hour. The solvents were removed under reduced pressure and the title compound was

purified by column chromatography ( $\text{SiO}_2$ ) eluting with a linear gradient (0 $\rightarrow$ 30% ethyl acetate-petrol) 16 mg (60%); LCMS (PS-A)  $R_t$  2.85 min;  $m/z$   $[\text{M}+\text{H}]^+$  266.

**8D. Preparation of Compounds of the Formula (I) from [4-(5-Methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-acetonitrile**

5 (i) The product of Example 8B can be reacted with benzaldehyde under the conditions described in Example 2 to give 2-[4-(5-methyl-1-(tetrahydro-pyran-2-yl)-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-3-phenyl-propionitrile which can be deprotected by removal of the 1-tetrahydropyranyl group under the conditions set out in Example 8C to give 2-[4-(5-methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-3-phenyl-propionitrile.

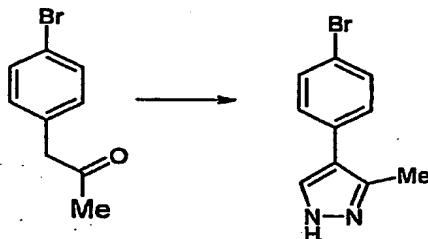
10 2-[4-(5-Methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-3-phenyl-propionitrile or its 1-tetrahydropyranyl derivative can be reduced according to the method of Example 6 (and thereafter where necessary deprotected according to the method of Example 8C) to give 2-[4-(5-methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-3-phenyl-propylamine.

15 The product of Example 8B can also be reacted with benzyl magnesium bromide or phenyl magnesium bromide under the Grignard reaction conditions described in Example 5 to give (following deprotection by the method of Example 8C) 1-benzyl-2-[4-(5-methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-ethylamine and 2-[4-(5-methyl-3-trifluoromethyl-1H-pyrazol-4-yl)-phenyl]-1-phenyl-ethylamine respectively.

**EXAMPLE 9**

**Construction of Pyrazole Ring System**

**9A. Synthesis of 4-(4-Bromo-phenyl)-3-methyl-1H-pyrazole**



To 4-bromophenylacetone (5.0 g, 23.5 mmol, 1.0 equiv) (Acros Organics 34216) was added N,N-dimethylformamide dimethyl acetal (11.3 ml, 84.6 mmol, 3.6 equiv) and the mixture was heated to 90 °C for 6 hours. The solvents were removed and the resulting gum was dissolved in ethanol (235 ml) with additional heating. Hydrazine hydrate (1.37 ml, 28.2 mmol, 1.2 equiv) was added and the mixture was heated to reflux for 15 hours. The solvents were removed under reduced pressure and the solid was triturated with dichloromethane to afford the title compound, 2.24 g (40%); LCMS (PS-A)  $R_t$  2.87 min;  $m/z$  [M+H]<sup>+</sup> 238. Further material could be isolated from the mother liquor.

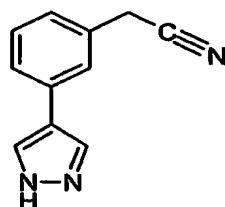
**9B. Conversion of 4-(4-Bromo-phenyl)-3-methyl-1H-pyrazole to compounds of the Formula (I)**

(i) 4-(4-Bromo-phenyl)-3-methyl-1H-pyrazole can be protected at the 1-position of the pyrazole ring by formation of the tetrahydropyranyl (THP) derivative by following the procedure set out in Example 8A. A Grignard reagent can then be prepared from the bromo-phenyl moiety by treating the protected derivative with magnesium in an ether solvent in standard fashion (see J. March, *Advanced Organic Chemistry*, 4<sup>th</sup> Edition, 1992, John Wiley, New York, pages 622-625). The Grignard reagent can be reacted with nitrostyrene (the nitrostyrene having been prepared by a standard method such as the method described in *Organic Syntheses*, Collective Volume 1, page 413) and the resulting nitroethyl compound reduced to give 2-{4-[3-methyl-1-(tetrahydro-pyran-2-yl)-1H-pyrazol-4-yl]-phenyl}-2-phenyl-ethylamine. Removal of the tetrahydropyranyl group using the method of Example 8C gives 2-{4-[3-methyl-1H-pyrazol-4-yl]-phenyl}-2-phenyl-ethylamine.

(ii) The bromo-compound of Example 9A can be converted into compounds of the formula (I) in which the group A contains a nitrogen atom which is attached to the group E. The introduction of a nitrogen containing entity can be accomplished by reaction of the compound of Example 9A with [3-(4-chlorophenylamino)-5-propyl]-methyl-carbamic acid tert-butyl ester under palladium catalysed amination conditions of the type described in *Organic Letters*, 2002, vol. 4, No. 17, pp2885-2888, followed by removal of the *t*-butyloxycarbonyl protecting group by standard methods.

**EXAMPLE 10**

10 [3-(1H-Pyrazol-4-yl)-phenyl]-acetonitrile



By following the procedure set out in Example 1 but using 3-bromophenyl-acetonitrile instead of 2-(4-chlorophenyl)-2-phenylethylamine, the title compound was obtained. LCMS (PS-A) 2.35 min;  $m/z$   $[M+H]^+$  184.

15 3-(1H-Pyrazol-4-yl)-phenyl-acetonitrile can be used as an intermediate in the preparation of compounds of the formula (I), for example by means of an aldehyde condensation reaction as described in Example 2 or a Grignard reaction as described in Example 5.

**BIOLOGICAL ACTIVITY**

20 EXAMPLE 11

Measurement of PKA Kinase Inhibitory Activity ( $IC_{50}$ )

Compounds of the invention can be tested for PKA inhibitory activity using the PKA catalytic domain from Upstate Biotechnology (#14-440) and the 9 residue PKA specific peptide (GRTGRRNSI), also from Upstate Biotechnology (#12-257), as the

substrate. A final concentration of 1 nM enzyme is used in a buffer that includes 20 mM MOPS pH 7.2, 40  $\mu$ M ATP/ $\gamma^{33}$ P-ATP and 50 mM substrate. Compounds are added in dimethylsulphoxide (DMSO) solution to a final DMSO concentration of 2.5%. The reaction is allowed to proceed for 20 minutes before addition of excess 5 orthophosphoric acid to quench activity. Unincorporated  $\gamma^{33}$ P-ATP is then separated from phosphorylated proteins on a Millipore MAPH filter plate. The plates are washed, scintillant is added and the plates are then subjected to counting on a Packard Topcount.

10 The % inhibition of the PKA activity is calculated and plotted in order to determine the concentration of test compound required to inhibit 50% of the PKB activity (IC<sub>50</sub>).

The compounds of Examples 1 and 4 have IC<sub>50</sub> values of less than 1  $\mu$ M whereas the compounds of Examples 5 and 7 have IC<sub>50</sub> values of less than 15  $\mu$ M.

#### EXAMPLE 12

15 **Measurement of PKB Kinase Inhibitory Activity (IC<sub>50</sub>)**

The inhibition of protein kinase B (PKB) activity by compounds can be determined determined essentially as described by Andjelkovic *et al.* (Mol. Cell. Biol. 19, 5061-5072 (1999)) but using a fusion protein described as PKB-PIF and described in full by Yang *et al* (Nature Structural Biology 9, 940 – 944 (2002)). The protein 20 is purified and activated with PDK1 as described by Yang *et al*. The peptide AKTide-2T (H-A-R-K-R-E-R-T-Y-S-F-G-H-H-A-OH) obtained from Calbiochem (#123900) is used as a substrate. A final concentration of 0.6 nM enzyme is used in a buffer that includes 20 mM MOPS pH 7.2, 30  $\mu$ M ATP/ $\gamma^{33}$ P-ATP and 25  $\mu$ M substrate. Compounds are added in DMSO solution to a final DMSO concentration 25 of 2.5%. The reaction is allowed to proceed for 20 minutes before addition of excess orthophosphoric acid to quench activity. The reaction mixture is transferred to a phosphocellulose filter plate where the peptide binds and the unused ATP is washed away. After washing, scintillant is added and the incorporated activity measured by scintillation counting.

The % inhibition of the PKa activity is calculated and plotted in order to determine the concentration of test compound required to inhibit 50% of the PKB activity (IC<sub>50</sub>).

Following the protocol described above, the IC<sub>50</sub> values of the compounds of

5 Examples 1 and 4 have been found to be less than 1  $\mu$ M whilst the compounds of Examples 2, 3, 5 and 6 each have IC<sub>50</sub> values of less than 5  $\mu$ M.

#### PHARMACEUTICAL FORMULATIONS

##### EXAMPLE 13

###### (i) Tablet Formulation

10 A tablet composition containing a compound of the formula (I) is prepared by mixing 50 mg of the compound with 197mg of lactose (BP) as diluent, and 3 mg magnesium stearate as a lubricant and compressing to form a tablet in known manner.

###### (ii) Capsule Formulation

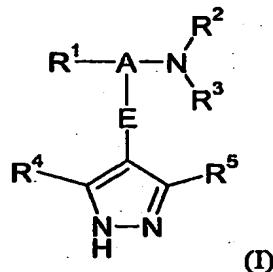
15 A capsule formulation is prepared by mixing 100mg of a compound of the formula (I) with 100mg lactose and filling the resulting mixture into standard opaque hard gelatin capsules.

##### Equivalents

The foregoing examples are presented for the purpose of illustrating the invention  
20 and should not be construed as imposing any limitation on the scope of the invention. It will readily be apparent that numerous modifications and alterations may be made to the specific embodiments of the invention described above and illustrated in the examples without departing from the principles underlying the invention. All such modifications and alterations are intended to be embraced by  
25 this application.

CLAIMS

## 1. A compound of the formula (I):



wherein A is a saturated hydrocarbon linker group containing from 1 to 7 carbon atoms, the linker group having a maximum chain length of 5 atoms extending between R<sup>1</sup> and NR<sup>2</sup>R<sup>3</sup> and a maximum chain length of 4 atoms extending between E and NR<sup>2</sup>R<sup>3</sup>, wherein one of the carbon atoms in the linker group may optionally be replaced by an oxygen or nitrogen atom; and wherein the carbon atoms of the linker group A may optionally bear one or more substituents selected from fluorine and hydroxy, provided that the hydroxy group is not located at a carbon atom  $\alpha$  with respect to the NR<sup>2</sup>R<sup>3</sup> group;

E is a monocyclic or bicyclic carbocyclic or heterocyclic group;

R<sup>1</sup> is an aryl or heteroaryl group;

R<sup>2</sup> and R<sup>3</sup> are independently selected from hydrogen, C<sub>1-4</sub> hydrocarbyl and C<sub>1-4</sub> acyl;

or R<sup>2</sup> and R<sup>3</sup> together with the nitrogen atom to which they are attached form a saturated monocyclic heterocyclic group having 4-7 ring members and optionally containing a second heteroatom ring member selected from O and N;

or one of R<sup>2</sup> and R<sup>3</sup> together with the nitrogen atom to which they are attached and one or more atoms from the linker group A form a saturated monocyclic heterocyclic group having 4-7 ring members and

optionally containing a second heteroatom ring member selected from O and N;

or  $NR^2R^3$  and the carbon atom of linker group A to which it is attached together form a cyano group;

5         $R^4$  is selected from hydrogen, halogen,  $C_{1-5}$  saturated hydrocarbyl, cyano and  $CF_3$ ; and

$R^5$  is selected from selected from hydrogen,  $C_{1-5}$  saturated hydrocarbyl, cyano,  $CONH_2$ ,  $CF_3$ ,  $NH_2$ ,  $NHCOR^9$  or  $NHCONHR^9$ ;

10         $R^9$  is phenyl or benzyl each optionally substituted by one or substituents selected from halogen, hydroxy, trifluoromethyl, cyano, nitro, carboxy, amino, mono- or di- $C_{1-4}$  hydrocarbylamino; a group  $R^a-R^b$  wherein  $R^a$  is a bond, O, CO,  $X^1C(X^2)$ ,  $C(X^2)X^1$ ,  $X^1C(X^2)X^1$ , S, SO,  $SO_2$ ,  $NR^c$ ,  $SO_2NR^c$  or  $NR^cSO_2$ ; and  $R^b$  is selected from hydrogen, heterocyclic groups having from 3 to 12 ring members, and a  $C_{1-8}$  hydrocarbyl group optionally substituted by one or more substituents selected from hydroxy, oxo, halogen, cyano, nitro, carboxy, amino, mono- or di- $C_{1-4}$  hydrocarbylamino, carbocyclic and heterocyclic groups having from 3 to 12 ring members and wherein one or more carbon atoms of the  $C_{1-8}$  hydrocarbyl group may optionally be replaced by O, S, SO,  $SO_2$ ,  $NR^c$ ,  $X^1C(X^2)$ ,  $C(X^2)X^1$  or  $X^1C(X^2)X^1$ ;

20         $R^c$  is selected from hydrogen and  $C_{1-4}$  hydrocarbyl; and  
 $X^1$  is O, S or  $NR^c$  and  $X^2$  is =O, =S or = $NR^c$ .

2. A compound according to claim 1 wherein the linker group A has a maximum chain length of 3 atoms (more preferably 1 or 2 atoms, and most preferably 2 atoms) extending between  $R^1$  and  $NR^2R^3$ .
- 25        3. A compound according to claim 1 or claim 2 wherein the linker group A has a maximum chain length of 3 atoms extending between E and  $NR^2R^3$ .
- 30        4. A compound according to claim 3 wherein the linker group A has a chain length of 2 or 3 atoms extending between  $R^1$  and  $NR^2R^3$  and a chain length of 2 or 3 atoms extending between E and  $NR^2R^3$ .

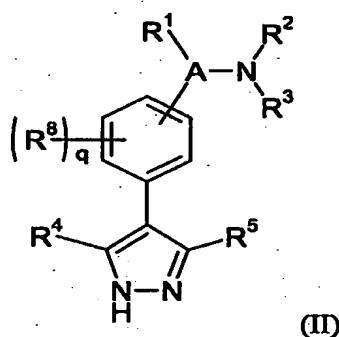
5. A compound according to any one of the preceding claims wherein the linker group atom linked directly to the group E is a carbon atom and the linker group A has an all-carbon skeleton.
6. A compound according to any one of the preceding claims wherein the moiety  $R^1-A-NR^2R^3$  is represented by the formula  $R^1-(G)_k-(CH_2)_m-X-(CH_2)_n-(CR^6R^7)_p-NR^2R^3$  wherein G is NH, NMe or O; X is attached to the group E and is selected from  $(CH_2)_j-CH$ ,  $(CH_2)_j-N$  and  $(NH)_j-CH$ ; j is 0 or 1, k is 0 or 1, m is 0 or 1, n is 0, 1, 2, or 3 and p is 0 or 1, and the sum of j, k, m, n and p does not exceed 4; and  $R^6$  and  $R^7$  are the same or different and are selected from methyl and ethyl, or  $CR^6R^7$  forms a cyclopropyl group.
7. A compound according to claim 6 wherein k is 0, m is 0 or 1, n is 0, 1, 2 or 3 and p is 0.
8. A compound according to claim 6 wherein k is 0, m is 0 or 1, n is 0, 1 or 2 and p is 1.
- 15 9. A compound according to claim 6 wherein X is  $(CH_2)_j-CH$ , k is 1, m is 0, n is 0, 1, 2 or 3 and p is 0.
10. A compound according to claim 6 wherein X is  $(CH_2)_j-CH$ , k is 1, m is 0, n is 0, 1 or 2 and p is 1.
11. A compound according to any one of claims 6, 9 and 10 wherein j is 0.
- 20 12. A compound according to any one of claims 6, 9 and 10 wherein j is 1.
13. A compound according to any one of claims 6, 8 and 10 wherein  $CR^6R^7$  is  $C(CH_3)_2$ .
14. A compound according to claim 1 wherein  $R^1-A(E)-NR^2R^3$  is a group selected from the groups A1 to A9 set out in Table 1 herein.

15. A compound according to claim 14 wherein  $R^1-A(E)-NR^2R^3$  is selected from groups A1, A2 and A3 in Table 1.
16. A compound according to any one of the preceding claims wherein E is a monocyclic group.
- 5 17. A compound according to any one of the preceding claims wherein E is an aryl or heteroaryl group.
18. A compound according to claim 17 wherein E is selected from optionally substituted phenyl, thiophene, furan, pyrimidine and pyridine groups.
19. A compound according to claim 18 wherein E is a phenyl group.
- 10 20. A compound according to any one of claims 1 to 16 wherein E is a non-aromatic monocyclic group selected from cycloalkanes such as cyclohexane and cyclopentane, and nitrogen-containing rings such as piperazine and piperazone.
21. A compound according to any one of the preceding claims wherein the group A and the pyrazole group are attached to the group E in a *meta* or *para* relative orientation; i.e. A and the pyrazole group are not attached to adjacent ring members of the group E.
- 15 22. A compound according to claim 21 wherein E is selected from 1,4-phenylene, 1,3-phenylene, 2,5-pyridylene and 2,4-pyridylene, 1,4-piperazinyl, and 1,4-piperazonyl.
- 20 23. A compound according to any one of the preceding claims wherein E is unsubstituted or has up to 4 substituents  $R^8$  selected from hydroxy, oxo (when E is non-aromatic), chlorine, bromine, trifluoromethyl, cyano,  $C_{1-4}$  hydrocarbyloxy and  $C_{1-4}$  hydrocarbyl optionally substituted by  $C_{1-2}$  alkoxy or hydroxy.
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24. A compound according to claim 23 wherein E has 0-3 substituents, more preferably 0-2 substituents, for example 0 or 1 substituent.

25. A compound according to claim 24 wherein E is unsubstituted.

26. A compound according to claim 23 having the formula (II):



wherein the group A is attached to the *meta* or *para* position of the benzene ring and q is 0-4.

27. A compound according to claim 26 wherein q is 0, 1 or 2, preferably 0 or 1 and most preferably 0.

10 28. A compound according to any one of the preceding claims wherein R<sup>1</sup> is selected from phenyl, naphthyl, thienyl, furan, pyrimidine and pyridine.

29. A compound according to claim 28 wherein R<sup>1</sup> is phenyl.

30. A compound according to any one of the preceding claims wherein R<sup>1</sup> is unsubstituted or is substituted by up to 5 substituents selected from hydroxy, C<sub>1-4</sub> acyloxy, fluorine, chlorine, bromine, trifluoromethyl, cyano, C<sub>1-4</sub> hydrocarbyloxy and C<sub>1-4</sub> hydrocarbyl optionally substituted by C<sub>1-2</sub> alkoxy or hydroxy.

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31. A compound according to claim 30 wherein R<sup>1</sup> is unsubstituted or is substituted by 0, 1, 2, 3 or 4 substituents, preferably 0, 1, 2 or 3, and more preferably 0, 1 or 2 substituents.
- 5 32. A compound according to claim 31 wherein the group R<sup>1</sup> has one or two substituents selected from fluorine, chlorine, trifluoromethyl, methyl and methoxy.
33. A compound according to claim 32 wherein R<sup>1</sup> is a mono-chlorophenyl or dichlorophenyl group.
- 10 34. A compound according to any one of the preceding claims wherein R<sup>4</sup> is selected from hydrogen and methyl.
- 15 35. A compound according to any one of the preceding claims wherein R<sup>5</sup> is selected from hydrogen, fluorine, chlorine, bromine, methyl, ethyl, hydroxyethyl, methoxymethyl, cyano, CF<sub>3</sub>, NH<sub>2</sub>, NHCOR<sup>9a</sup> and NHCONHR<sup>9a</sup> where R<sup>9a</sup> is phenyl or benzyl optionally substituted by hydroxy, C<sub>1-4</sub> acyloxy, fluorine, chlorine, bromine, trifluoromethyl, cyano, C<sub>1-4</sub> hydrocarbyloxy and C<sub>1-4</sub> hydrocarbyl optionally substituted by C<sub>1-2</sub> alkoxy or hydroxy.
- 20 36. A compound according to any one of the preceding claims having a molecular weight no greater than 1000, more usually less than 750, for example less than 700, or less than 650, or less than 600, or less than 550.
37. A compound according to claim 36 wherein the molecular weight is less than 525 and, for example, is 500 or less.
- 25 38. A compound of the formula (I) as defined in any of the examples herein.
39. A compound according to any one of the preceding claims in the form of a salt, solvate (such as a hydrate), ester or N-oxide.

40. A compound as defined in any one of claims 1 to 39 for use in the prophylaxis or treatment of a disease state or condition mediated by protein kinase B.
41. The use of a compound as defined in any one of claims 1 to 39 for the manufacture of a medicament for the prophylaxis or treatment of a disease state or condition mediated by protein kinase B.
42. A method for the prophylaxis or treatment of a disease state or condition mediated by protein kinase B, which method comprises administering to a subject in need thereof a compound as defined in any one of claims 1 to 39.
43. A method for treating a disease or condition comprising or arising from abnormal cell growth in a mammal, which method comprises administering to the mammal a compound as defined in any one of claims 1 to 39 in an amount effective in inhibiting abnormal cell growth.
44. A method for treating a disease or condition comprising or arising from abnormal cell growth in a mammal, the method comprising administering to the mammal a compound as defined in any one of claims 1 to 39 in an amount effective to inhibit PKB activity.
45. A method of inhibiting a protein kinase B, which method comprises contacting the kinase with a kinase-inhibiting compound as defined in any one of claims 1 to 39.
46. A method of modulating a cellular process by inhibiting the activity of a protein kinase B using a compound as defined in any one of claims 1 to 39.
47. A method for treating an immune disorder in a mammal, the method comprising administering to the mammal a compound as defined in any one of claims 1 to 39 in an amount effective to inhibit PKB activity.

48. A compound as defined in any one of claims 1 to 39 for use in the prophylaxis or treatment of a disease state or condition mediated by protein kinase A.

49. The use of a compound as defined in any one of claims 1 to 39 for the manufacture of a medicament for the prophylaxis or treatment of a disease state or condition mediated by protein kinase A.

50. The use of a compound of the formula (I) as defined in any one of claims 1 to 39 for the manufacture of a medicament for the prophylaxis or treatment of a disease state or condition arising from abnormal cell growth.

10 51. The use of a compound of the formula (I) as defined in any one of claims 1 to 39 for the manufacture of a medicament for the prophylaxis or treatment of a disease in which there is a disorder of proliferation, apoptosis or differentiation.

15 52. A method for the prophylaxis or treatment of a disease state or condition mediated by protein kinase A, which method comprises administering to a subject in need thereof a compound as defined in any one of claims 1 to 39.

53. A method for treating a disease or condition comprising or arising from abnormal cell growth in a mammal, the method comprising administering to the mammal a compound as defined in any one of claims 1 to 39 in an amount effective to inhibit PKA.

20 54. A method of inhibiting a protein kinase A, which method comprises contacting the kinase with a kinase-inhibiting compound as defined in any one of claims 1 to 39.

55. A method of modulating a cellular process by inhibiting the activity of a protein kinase A using a compound as defined in any one of claims 1 to 39.

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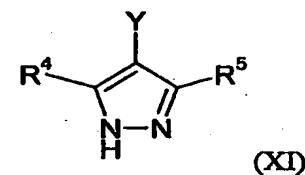
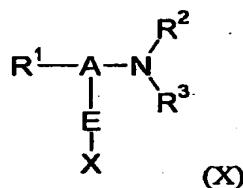
56. A method for treating an immune disorder in a mammal, the method comprising administering to the mammal a compound as defined in any one of claims 1 to 39 in an amount effective to inhibit PKA activity.

57. A method of inducing apoptosis in a cancer cell, which method comprises contacting the cancer cell with a compound as defined in any one of claims 1 to 39.

58. A pharmaceutical composition comprising a novel compound as defined in any one of claims 1 to 39 and a pharmaceutically acceptable carrier.

59. A compound as defined in any one of claims 1 to 39 for use in medicine.

10 60. A process for the preparation of a compound of the formula (I) as defined in any one of claims 1 to 39, which process comprises the reaction of a compound of the formula (X) with a compound of the formula (XI) or an N-protected derivative thereof:



15 wherein A, E, and R<sup>1</sup> to R<sup>5</sup> are as defined in any one of the preceding claims, one of the groups X and Y is selected from chlorine, bromine, iodine and trifluoromethanesulphonate, and the other one of the groups X and Y is a boronate residue, for example a boronate ester or boronic acid residue, in the presence of a palladium catalyst and a base.